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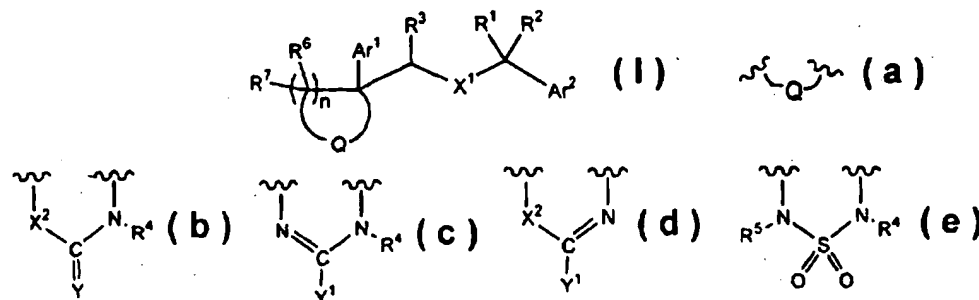
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(54) Title: SELECTIVE NEUROKININ ANTAGONISTS





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SELECTIVE NEUROKININ ANTAGONISTS

10 BACKGROUND OF THE INVENTION

The present invention relates to a genus of substituted cyclic ureas and derivatives thereof useful as antagonists of tachykinin receptors, in particular as antagonists of the neuropeptides neurokinin-1 receptor (NK₁).

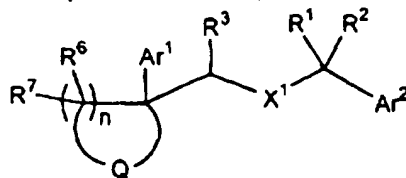
15 Neurokinin receptors are found in the nervous system and the circulatory system and peripheral tissues of mammals, and therefore are involved in a variety of biological processes. Neurokinin receptor antagonists are consequently expected to be useful in the treatment or prevention of various mammalian disease states, for example respiratory diseases such as chronic lung disease, bronchitis, pneumonia, asthma, allergy, cough, bronchospasm; inflammatory diseases such as arthritis and psoriasis; skin disorders such as atopic dermatitis and contact dermatitis; 20 ophthalmological disorders such as retinitis, ocular hypertension and cataracts; addictions such as alcohol dependence and psychoactive substance abuse; stress related disorders such as post traumatic stress disorder; obsessive/ compulsive disorders; eating disorders such as bulimia, anorexia nervosa and binge eating disorders; mania; premenstrual syndrome; central nervous system conditions such 25 as anxiety, general anxiety disorder, panic disorder, phobias, bipolar disorders, migraine, epilepsy, nociception, emesis, depression, psychosis, schizophrenia, Alzheimer's disease, AIDs related dementia and Towne's disease; gastrointestinal disorders such as Crohn's disease and colitis; nausea; bladder disorders; atherosclerosis; fibrosing disorders; obesity; Type II diabetes; pain related disorders 30 such as neuropathic pain, post-operative pain, headache and chronic pain syndromes; and genitourinary disorders such as interstitial cystitis and urinary incontinence.

In particular, NK₁ receptors have been reported to be involved in microvascular 35 leakage and mucus secretion, making NK₁ receptor antagonists especially useful in the treatment and prevention of asthma, emesis, nausea, depression, anxiety, cough, pain and migraine.

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SUMMARY OF THE INVENTION

Compounds of the present invention are represented by the formula I



or a pharmaceutically acceptable salt thereof, wherein

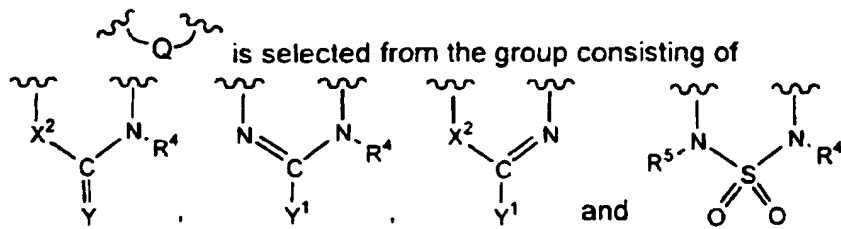
- 5 Ar¹ and Ar² are independently selected from the group consisting of R¹⁷-heteroaryl and

X¹ is -O-, -S-, -SO-, -SO₂-, -NR¹²-, -N(COR¹²)- or -N(SO₂R¹⁵)-;

- 10 R¹, R², R³ and R⁷ are each independently selected from the group consisting of H, C₁-C₆ alkyl, hydroxy(C₁-C₃)alkyl, C₃-C₈ cycloalkyl, -CH₂F, -CHF₂ and -CF₃; or R¹ and R², together with the carbon to which they are attached, form a C₃-C₆ alkylene ring; or, when X¹ is -O-, -S- or -NR¹²-, R¹ and R² together are =O;

each R⁶ is independently selected from H, C₁-C₆ alkyl, -OR¹³ or -SR¹²;n is 1-4, if n is greater than 1, then R⁶ and R⁷ can be the same or different on

- 15 each carbon;

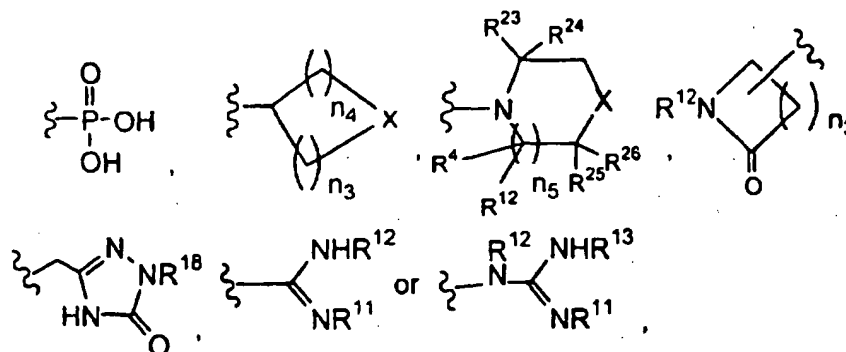
X² is -O-, -S- or -NR⁵-;Y is =O, =S or =NR¹¹;

- 20 Y¹ is H, C₁-C₆ alkyl, -NR¹⁷R¹³, -SCH₃, R¹⁹-aryl(CH₂)_{n6}-, R¹⁹-heteroaryl-(CH₂)_{n6}-, -(CH₂)_{n6}-heterocycloalkyl, -(C₁-C₃)alkyl-NH-C(O)O(C₁-C₆)alkyl or -NHC(O)R¹⁵;

R⁵ is H or -(CH₂)_{n1}-G, wherein n₁ is 0-5, G is H, -CF₃, -CHF₂, -CH₂F, -OH, -O-(C₁-C₆ alkyl), -SO₂R¹³, -O-(C₃-C₈ cycloalkyl), -NR¹³R¹⁴, -SO₂NR¹³R¹⁴, -NR¹³SO₂R¹⁵, -NR¹³COR¹², -NR¹²(CONR¹³R¹⁴), -CONR¹³R¹⁴, -COOR¹², C₃-C₈ cycloalkyl, R¹⁹-aryl, R¹⁹-

- 25 heteroaryl,

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or when n_1 is 0, R^5 can also be $-C(O)R^{13}$ or $-C(S)R^{13}$; provided that G is not H when $n_1=0$;

X is $-NR^{20}$ -, $-N(CONR^{13}R^{14})$ -, $-N(CO_2R^{13})$ -, $-N(SO_2R^{15})$ -, $-N(COR^{12})$ -,

5 $N(SO_2NHR^{13})$ -, $-O$ -, $-S$ -, $-SO$ -, $-SO_2$ -, $-CF_2$ -, $-CH_2$ - or $-CR^{12}F$ -;

R^8 , R^9 and R^{10} are independently selected from the group consisting of H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, $-OR^{12}$, halogen, $-CN$, $-NO_2$, $-CF_3$, $-CHF_2$, $-CH_2F$, $-OCF_3$, $-OCHF_2$, $-OCH_2F$, $-COOR^{12}$, $-CONR^{21}R^{22}$, $-NR^{21}COR^{12}$, $-NR^{21}CO_2R^{15}$, $-NR^{21}CONR^{21}R^{22}$, $-NR^{21}SO_2R^{15}$, $-NR^{21}R^{22}$, $-SO_2NR^{21}R^{22}$, $-S(O)_{n_5}R^{15}$, R^{16} -aryl and R^{16} -heteroaryl;

10 R^{11} is H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, $-NO_2$, $-CN$, OH , $-OR^{12}$, $-O(CH_2)_{n_6}R^{12}$, $-(C_1-C_3)alkyl-C(O)NHR^{12}$, R^{16} -aryl $(CH_2)_{n_6}$ - or R^{16} -heteroaryl $(CH_2)_{n_6}$;

R^4 and R^{12} are each independently selected from the group consisting of H, C_1 - C_6 alkyl and C_3 - C_8 cycloalkyl;

15 R^{13} and R^{14} are independently selected from the group consisting of H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, R^{16} -aryl $(CH_2)_{n_6}$ - or R^{16} -heteroaryl $(CH_2)_{n_6}$;; or R^{13} and R^{14} together are C_3 - C_6 alkylene and with the nitrogen to which they are attached form a 4-7 membered ring, or one of the carbon atoms in the alkylene chain formed by R^{13} and R^{14} is replaced by a heteroatom selected from the group consisting of $-O$ -, $-S$ - and $-NR^{12}$;;

20 R^{15} is C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl or $-CF_3$;

R^{16} is 1 to 3 substituents independently selected from the group consisting of H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, C_1 - C_6 alkoxy, halogen and $-CF_3$;

R^{17} is H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, $-COOR^{12}$, $-CONR^{21}R^{22}$, $-NR^{21}R^{22}$, $-NR^{21}COR^{12}$, $-NR^{21}CO_2R^{12}$, $-NR^{21}CONR^{21}R^{22}$, $-NR^{21}SO_2R^{15}$ or $-S(O)_{n_5}R^{15}$;

25 R^{18} is H, C_1 - C_6 alkyl or $-P(O)(OH)_2$;

R^{19} is 1 to 3 substituents independently selected from the group consisting of H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, OH , halogen, $-CN$, $-NO_2$, $-CF_3$, $-CHF_2$, $-CH_2F$, $-OCF_3$, $-OCHF_2$, $-OCH_2F$, $-O(C_1-C_6 alkyl)$, $-O(C_3-C_8 cycloalkyl)$, $-COOR^{12}$, $-CONR^{21}R^{22}$, $-NR^{21}R^{22}$, $-NR^{21}COR^{12}$, $-NR^{21}CO_2R^{12}$, $-NR^{21}CONR^{21}R^{22}$, $-NR^{21}SO_2R^{15}$ and $-S(O)_{n_5}R^{15}$;

30 R^{20} is H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl or $-(CH_2)_{n_6}$ -heterocycloalkyl;

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R^{21} and R^{22} are independently selected from the group consisting of H, C_1 - C_6 alkyl, C_3 - C_6 cycloalkyl and benzyl; or R^{21} and R^{22} together are C_3 - C_6 alkylene and with the nitrogen to which they are attached form a 4-7 membered ring, or one of the carbon atoms in the alkylene chain formed by R^{21} and R^{22} is replaced by a
 5 heteroatom selected from the group consisting of -O-, -S- and -NR¹²-;

R^{23} , R^{24} , R^{25} and R^{26} are H, C_1 - C_6 alkyl and can be together as =O; when $n_3 = 0$, and R^{25} and $R^{26} = H$, X is not O, N, S;

n_3 and n_4 are independently 1-5, provided that the sum of n_3 and n_4 is 2-6;

n_5 is independently 0-2;

10 n_6 is independently 0-3; and

q and r are independently 1 or 2.

Preferred are compounds of formula I wherein R^4 and R^7 are each H. Also preferred are compounds of formula I wherein R^1 and R^3 are each H. Also preferred are compounds of formula I wherein R^1 , R^3 , R^4 and R^7 are each H. R^6 is preferably H
 15 or -OH. Preferably, X^1 is -O- or -NR¹²-. Ar^1 and Ar^2 are each preferably R^8 , R^9 , R^{10} -phenyl, wherein R^8 , R^9 and R^{10} are independently selected. Y is preferably =O, and n is preferably 1 or 2. When Y is =O, X^2 is preferably -NR⁵-. More preferred are compounds of formula I wherein Q is -X²-C(=Y)-NR⁴- (i.e., the first structure shown in the definition of Q), R^1 , R^3 , R^4 and R^7 are each H; R^6 is H or -OH; X^1 is -O- or -NR¹²-
 20 ; Ar^1 and Ar^2 are each R^8 , R^9 , R^{10} -phenyl; Y is =O and X^2 is -NR⁵-; n is 1 or 2; R^5 is H or -(CH₂)_{n1}- G, G is not H when $n_1 = 0$; R^{19} -aryl or R^{19} -heteroaryl. Most preferred are compounds of formula I wherein R^5 is H.

This invention also relates to the use of a compound of formula I in the treatment of, for example, respiratory diseases such as chronic lung disease,
 25 bronchitis, pneumonia, asthma, allergy, cough, bronchospasm; inflammatory diseases such as arthritis and psoriasis; skin disorders such as atopic dermatitis and contact dermatitis; ophthalmological disorders such as retinitis, ocular hypertension and cataracts; addictions such as alcohol dependence and psychoactive substance abuse; stress related disorders such as post traumatic stress disorder; obsessive/
 30 compulsive disorders; eating disorders such as bulimia, anorexia nervosa and binge eating disorders; mania; premenstrual syndrome; central nervous system conditions such as anxiety, general anxiety disorder, panic disorder, phobias, bipolar disorders, migraine, epilepsy, nociception, emesis, depression, psychosis, schizophrenia, Alzheimer's disease, AIDs related dementia and Towne's disease; gastrointestinal
 35 disorders such as Crohn's disease and colitis; nausea; bladder disorders; atherosclerosis; fibrosing disorders; obesity; Type II diabetes; pain related disorders such as neuropathic pain, post-operative pain, headache and chronic pain syndromes; and genitourinary disorders such as interstitial cystitis and urinary

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incontinence. The treatment of mammals, both human and non-human, is contemplated.

Further, the invention relates to a method for antagonizing the effect of Substance P at its receptor site or for the blockade of neurokinin-1 receptors in a mammal, comprising administering an amount of a compound of formula I effective to antagonize the effect of Substance P at its receptor site in a mammal in need of such treatment.

In another aspect, the invention relates to a pharmaceutical composition comprising a compound of formula I in a pharmaceutically acceptable carrier. The invention also relates to the use of said pharmaceutical composition in the treatment of the mammalian disease states listed above.

The compounds of this invention can be combined with a selective serotonin reuptake inhibitor (SSRI) (i.e., the compounds of this invention can be combined with an SSRI in a pharmaceutical composition, or the compounds of this invention can be administered with an SSRI.

Numerous chemical substances are known to alter the synaptic availability of serotonin through their inhibition of presynaptic reaccumulation of neuronally released serotonin. Representative SSRIs include, without limitation; fluoxetine, fluvoxamine, paroxetine and sertraline, and pharmaceutically acceptable salts thereof. Other compounds can readily be evaluated to determine their ability to selectively inhibit serotonin reuptake.

In another aspect, the invention relates to a method of treating the above diseases and disorders comprising administering an effective amount of an NK1 antagonist of formula I in combination with an SSRI described above.

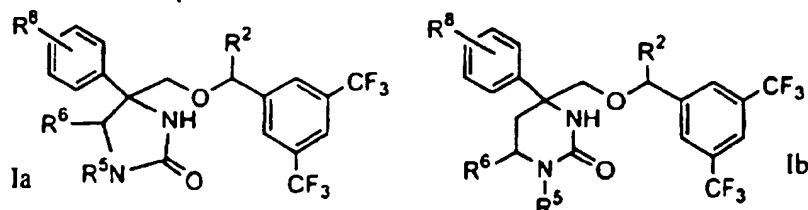
In another aspect, the invention relates to a method of treating the above diseases and disorders comprising administering an effective amount of an NK1 antagonist of formula I in combination with an SSRI selected from: fluoxetine, fluvoxamine, paroxetine and sertraline, and pharmaceutically acceptable salts thereof.

In another aspect, the invention relates to a method of treating emesis, depression, anxiety, and cough comprising administering an effective amount of an NK1 antagonist of formula I in combination with an SSRI described above.

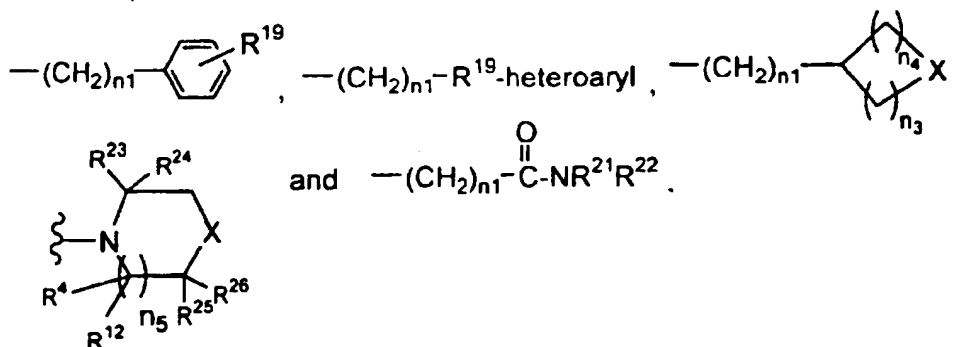
In the methods of this invention wherein a combination of an NK1 antagonist of this invention (compound of formula I) is administered with an SSRI described above, the compound of formula I and SSRI can be administered simultaneously, consecutively (one after the other within a relatively short period of time), or sequentially (first one and then the other over a period of time). In general, when the antagonists are administered consecutively or sequentially, the NK1 antagonist of this invention (compound of formula I) is administered first.

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Preferred are compounds of formula Ia and Ib

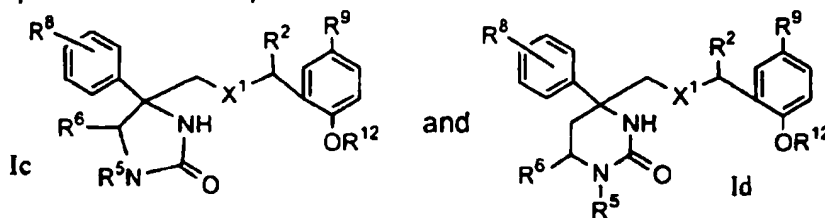


wherein R^8 is H or halogen; R^2 is H, $-CH_3$ or $-CH_2OH$; R^6 is H or $-OH$; and R^5 is selected from the group consisting of hydrogen and groups of the formula $-(CH_2)_{n1}-G$ as follows,



or $-(CH_2)_{n1}-G'$, wherein n_1 is 2-4 and G' is H, $-OH$, $-OCH_3$, $-OEt$, $-O(i-Pr)$, $-O$ -cyclopropyl, or $-CONR^{13}R^{14}$, wherein R^{13} and R^{14} are independently selected from the group consisting of H, $-CH_3$, Et, $i-Pr$, or cyclopropyl (Et is ethyl and $i-Pr$ is isopropyl).

Also preferred are compounds of the formula Ic and Id



wherein X^1 is $-O-$, $-NH-$, $-N(CH_3)-$ or $-N(COCH_3)-$; R^8 is H or halogen; R^2 is H, $-CH_3$ or $-CH_2OH$; R^9 is $-OCF_3$ or 5-(trifluoromethyl)-1H-tetrazol-1-yl; R^6 is H or $-OH$; R^{12} is $-CH_3$ or cyclopropyl; and R^5 is selected from the group consisting of hydrogen and groups of the formula $-(CH_2)_{n1}-G$ as shown above for structures Ia and Ib.

Preferred compounds of the invention are the compounds of examples 2, 61, 79a, 79b, 92, 93, 126, 127, 128, 129, 165a, 165b, 166a and 166b.

DETAILED DESCRIPTION

As used herein, the term "alkyl" means straight or branched alkyl chains. "Lower alkyl" refers to alkyl chains of 1-6 carbon atoms and, similarly, lower alkoxy refers to alkoxy chains of 1-6 carbon atoms.

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"Cycloalkyl" means cyclic alkyl groups having 3 to 8 carbon atoms.

"Aryl" means phenyl, naphthyl, indenyl, tetrahydronaphthyl, indanyl, anthracenyl or fluorenyl. R^{16} -aryl and R^{19} -aryl refer to such groups wherein substitutable ring carbon atoms have an R^{16} or an R^{19} substituent.

5 "Halogen" refers to fluoro, chloro, bromo or iodo atoms.

"Heteroaryl" refers to 5- to 10-membered single or benzofused aromatic rings comprising 1 to 4 heteroatoms independently selected from the group consisting of -O-, -S-, -N= and -NH-, provided that the rings do not include adjacent oxygen and/or sulfur atoms. Examples of single-ring heteroaryl groups are furanyl, imidazolyl, isoxazolyl, isothiazolyl, oxadiazolyl, oxazolyl, pyrazinyl, pyrazolyl, pyridyl, pyrimidyl, pyrrolyl, tetrazolyl, thiadiazolyl, thiazolyl, thienyl, triazolyl and triazinyl. Examples of benzofused heteroaryl groups are benzimidazolyl, benzofuranyl, benzo-thiophenyl, benzoxazolyl, indolyl and quinolyl. N-oxides of nitrogen-containing heteroaryl groups are also included. All positional isomers are contemplated, e.g., 2-pyridyl, 3-pyridyl, 10 and 4-pyridyl, and when R^4 or R^5 is heteroaryl, it can be joined to the nitrogen atom of the "-Q-" group either by a ring carbon or a ring nitrogen. R^{19} -heteroaryl refer to such groups wherein substitutable ring carbon atoms have an R^{19} substituent. When R^8 , R^9 or R^{10} is heteroaryl, it is preferably tetrazolyl substituted by H, C_1 - C_4 alkyl, C_3 - C_8 cycloalkyl, $-CF_3$, $-SO_2$ -(C_1 - C_6 alkyl) or $-OCF_3$.

20 "Heterocycloalkyl" refers to a 4- to 7-membered saturated ring comprising 1 to 3 heteroatoms independently selected from the group consisting of -O-, -S- and -NR²¹-, wherein R^{21} is H or C_1 - C_6 alkyl, and wherein the remaining ring members are carbon. Where a heterocyclic ring comprises more than one heteroatom, no rings are formed where there are adjacent oxygen atoms, adjacent sulfur atoms, or three consecutive heteroatoms. Examples of heterocyclic rings are tetrahydrofuranyl, pyrrolidinyl, tetrahydropyranyl, piperidinyl, morpholinyl, thiomorpholinyl and piperazinyl.

In the above definitions, wherein variables R^1 to R^{26} are said to be independently selected from a group of substituents, we mean that R^1 , R^2 , R^3 , etc., 30 are independently selected, but also that where an R^8 , for example, occurs more than once in a molecule, those occurrences are independently selected (e.g., if X^1 is $-NR^{12}$ - wherein R^{12} is hydrogen, G can be $-COOR^{12}$ wherein R^{12} is methyl). Similarly, R^8 , R^9 and R^{10} can be independently selected from a group of substituents, and where more than one R^8 , R^9 or R^{10} is other than hydrogen, the substituents are 35 independently selected; those skilled in the art will recognize that the size and nature of the substituent(s) will affect the number of substituents which can be present.

The "Q" groups are always joined to the rest of the molecule as shown, i.e., they are attached left-to-right, where $-X^2$ - or $-NR^5$ - is attached to the carbon to which

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R⁶ and R⁷ are attached, and -NR⁴- is always attached to the carbon to which Ar¹ is attached.

Compounds of formula I can have at least one asymmetrical carbon atom and all isomers, including diastereomers, enantiomers and rotational isomers are contemplated as being part of this invention. The invention includes d and l isomers in both pure form and in admixture, including racemic mixtures. Isomers can be prepared using conventional techniques, either by reacting optically pure or optically enriched starting materials or by separating isomers of a compound of formula I.

Those skilled in the art will appreciate that for some compounds of formula I, one isomer will show greater pharmacological activity than other isomers.

Compounds of the invention which have an amino group can form pharmaceutically acceptable salts with organic and inorganic acids. Examples of suitable acids for salt formation are hydrochloric, sulfuric, phosphoric, acetic, citric, oxalic, malonic, salicylic, malic, fumaric, tartaric, succinic, ascorbic, maleic, methanesulfonic and other mineral and carboxylic acids well known to those in the art. The salt is prepared by contacting the free base form with a sufficient amount of the desired acid to produce a salt. The free base form may be regenerated by treating the salt with a suitable dilute aqueous base solution such as dilute aqueous sodium bicarbonate. The free base form differs from its respective salt form somewhat in certain physical properties, such as solubility in polar solvents, but the salt is otherwise equivalent to its respective free base forms for purposes of the invention.

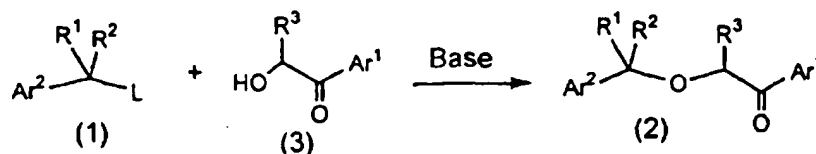
Certain compounds of the invention which are acidic (e.g., those compounds which possess a carboxyl group) form pharmaceutically acceptable salts with inorganic and organic bases. Examples of such salts are the sodium, potassium, calcium, aluminum, gold and silver salts. Also included are salts formed with pharmaceutically acceptable amines such as ammonia, alkyl amines, hydroxyalkylamines, N-methylglucamine and the like.

Compounds of formula I can be prepared using methods known to those skilled in the art. Typical procedures are described below, although the skilled artisan will recognize that other procedures may be applicable, and that the procedure may be suitably modified to prepare other compounds within the scope of formula I.

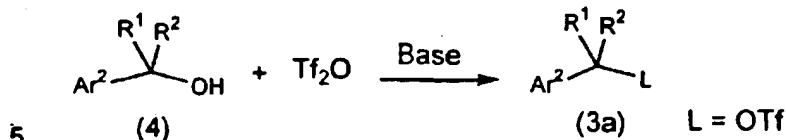
Method 1:

Compounds of formula I wherein X¹ is -O- or -S- and n is 1 can be prepared by the following method.

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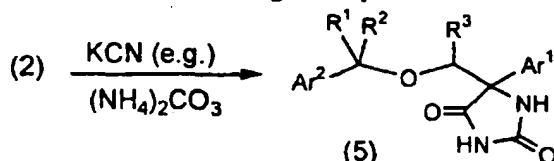
An alcohol (3), in which Ar¹ and R³ are as defined above is converted to the ketone (2), by reaction with an activated derivative (1) of the alcohol (4), in which Ar², R¹ and R² are as defined above.



This reaction is most favorable when R¹, R² and R³ are each H but, depending on the leaving group L, it may work effectively if either R¹, R² or R³ is C1-C6 alkyl. A leaving group, L, of choice is CF₃SO₂- (triflate) but others also suffice, such as Br or I. The base used may vary but is preferably one of the hindered non-nucleophilic kind, of which an example is 2,6-di-tert-butyl-4-methyl pyridine.

The alkylating agent (3a) in which L is triflate may be prepared from the alcohol-type starting material (4) using triflic anhydride and the same hindered, non-nucleophilic base as is used for the alkylation.

The ketone (2) may be used to prepare compounds in which n is 1, X² is -NR⁵- and Y is =O (5). Reaction of (2) with a metallic cyanide (e. g. KCN) and (NH₄)₂CO₃ results in the formation of the hydantoin, a process well known to those skilled in the art of organic synthesis:

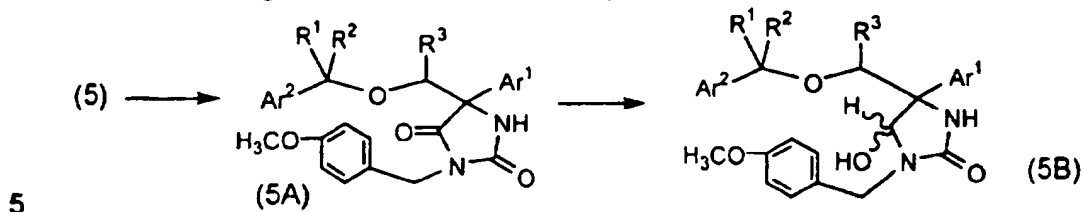


Selective reduction of the amide carbonyl and not the urea carbonyl may be accomplished by using a mixture of lithium aluminum hydride (LAH) and AlCl₃ as a preferred method although other methods are also available, such as the use of LAH in ether or THF at or above room temperature, up to the boiling point of the solution.

The reaction produces compounds of the invention in which R⁴ and R⁵ are both H. Introduction of a substituent, R⁵, may be performed relatively selectively although in some cases a second substituent R⁴ (where R⁴ = R⁵ in this case) may be introduced at the same time. Such substitution reaction at the nitrogen atoms of (5) may be accomplished using one of many sets of conditions used for such transformations, for example, use of an organic base, such as triethylamine or Hunig's Base (di-isopropyl ethylamine) and the appropriate alkylating agent, L-R⁵.

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Reaction of the hydantoin (5) with p-methoxybenzyl chloride in acetone in the presence of K_2CO_3 and a catalytic quantity of tetra-n-butylammonium iodide produces the derivative (5A) which can be readily reduced to a mixture of alcohols (5B) by use of mild reducing conditions. Suitable reagents are LAH in THF at 0-30°C for 1-6 hrs.

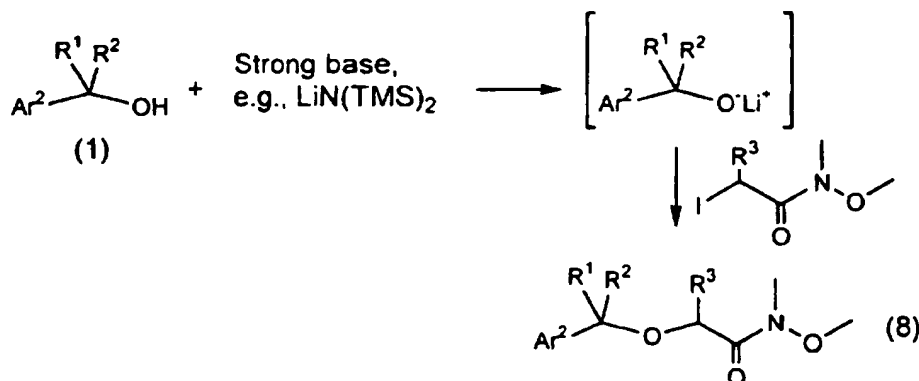


Subsequent removal of the PMB protecting group may be carried out using $Ce(NH_4)_2(NO_3)_6$ (CAN) in a neutral solvent, preferably a CH_3CN /water mixture. The mixture of chiral alcohols can frequently be separated and purified by chiral HPLC, preferably on one of the carbohydrate-based columns, such as one of the Daicel Chiralcel® or Chiralpak® series of columns.

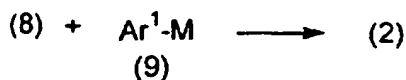
Method 2:

Compounds of formula I wherein X^1 is -O- or -S- and Y is =O or =S can be prepared by the following method.

15 The ketone (2) may also be made by the following sequence of reactions. The alcohol (1) may be converted to its alkoxide anion using a strong base, such as lithium bis(trimethyl silyl)amide or the like, followed by reaction, in an inert solvent, such as THF, with the N,O-dimethyl amide of the iodo-acid (7) to produce (8) which is known as a "Weinreb amide".

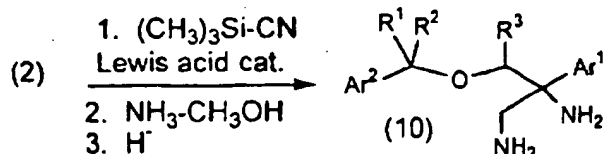


20 Addition of an organometallic derivative of Ar^1 (9) results in formation of the ketone (2). Suitable organometallic reagents include the Grignard (M is Mg) or lithium reagent. Suitable media for this reaction include neutral, non-reactive solvents such as ether or THF.



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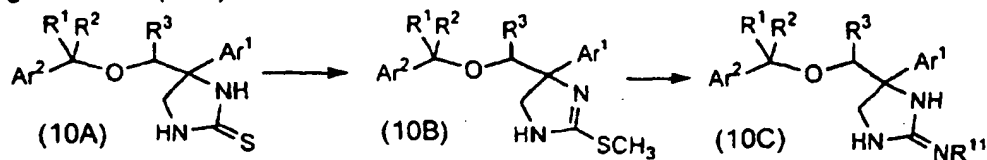
- 5 The ketone (2) may next be reacted with trimethylsilylcyanide in the presence of a Lewis acid catalyst, such as ZnI_2 , and subsequently treated with saturated $\text{NH}_3\text{-CH}_3\text{OH}$ at ambient temperature to yield an intermediate which may be reduced directly to diamine (10) using a powerful hydride reducing agent such as LAH in a neutral solvent such as THF.



- Reaction of the diamine (10) with a reagent known to introduce a carbonyl between two amines located in the correct position leads to the cyclic ureas (5) which are compounds of the invention. Examples of such reagents are COCl_2 , carbonyl diimidazole and methyl or ethyl chloroformate. Subsequent modification by introduction of R^4 and R^5 groups may be performed as described in Method 1.

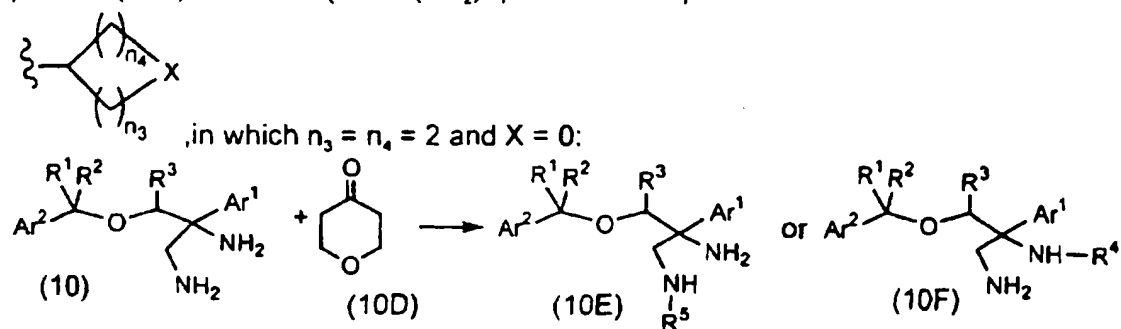
- The reaction described above can also be used to prepare compounds of the invention in which X^1 is $-\text{S}-$ by employing the thiol corresponding to the alcohol (4) shown above. In addition, reagents known to introduce the $-\text{C}(=\text{S})-$ function between two appropriately placed nitrogen atoms (such as thiocarbonyl diimidazole) may be used to prepare compounds in which Y is $=\text{S}$.

- A further use of compounds such as 10A is to introduce the guanidine functions into compounds of the invention. Reacting (10A) with CH_3I in a neutral solvent, such as CH_3CN , THF, or a mixture of the two, produces the S-methyl derivative (10B) which may then be reacted with an amine, $\text{R}^{11}\text{-NH}_2$, to produce guanidines (10C) or a tautomer.



- The diamine (10) is also a useful intermediate for preparation of products of the invention in which X^2 is $-\text{NR}^5-$. The group R^5 is introduced by use of an aldehyde or ketone precursor of R^5 by a process of reductive amination (otherwise known as reductive alkylation of the amine (10)). A proviso of this method is that the R^5 group may not contain a quaternary carbon atom next to the nitrogen atom. It also cannot be H, nor certain other of the definitions of R^5 , for example, when G is $-\text{OH}$, $-\text{SO}_2\text{R}^{13}$ or $-\text{NR}^{13}\text{R}^{14}$ etc., then n , cannot be 0. To describe the process, the starting material (10D) will be used. By reacting (10) with (10D) in a neutral solvent, such as 1,2-dichloroethane, in the presence of a suitable reducing agent (sodium triacetoxy-

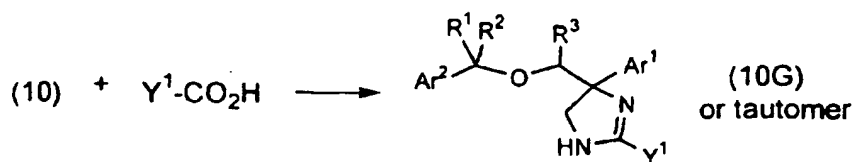
borohydride is particularly suitable for this reaction), and conventional work-up, a product (10E) is formed (R^5 is $-(CH_2)_{n_1}-G$ in which n_1 is 0 and G is



- 5 A variable amount of the isomeric structure (10F) (see above) may also form in this reaction, depending upon the reactants. It may be separated from (10E) by conventional chromatographic methods and reacted, as above, to yield compounds of the invention in which R^4 is the introduced substituent instead of R^5 .

- 10 Reactions with other precursors of the R^5 group are also possible, as will be evident to one skilled in the arts of organic and medicinal chemistry. Ring closing of (10E) may be carried out by direct cyclization to make many of the Q groups of the invention in which an R^5 group is present and where Y is $=O$ or $=S$, or compounds wherein Y is $=N-R^5$ can be made by the sequence of reactions described earlier for the synthesis of (10C).

- 15 A further use for the diamine (10) is in the preparation of compounds of the invention in which Y^1 is as defined above, except that it is not $-NH_2$, $-NHCH_3$ or $-SCH_3$. The diamine (10) may be heated with a carboxylic acid, Y^1-CO_2H , in a high boiling, neutral solvent, such as toluene to produce the amidines (10G) of the invention:

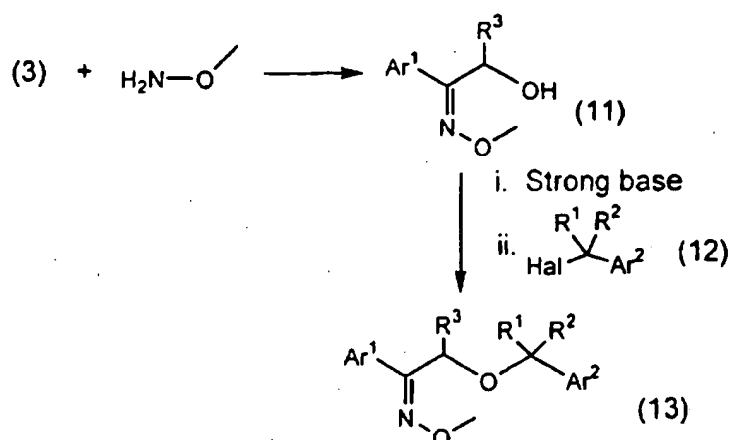


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Method 3:

- An alcohol (3) may be reacted with an O-substituted hydroxylamine derivative, preferably methoxyamine to yield the oxime derivative (11). Conversion of the oxime to the alkoxide may be performed using a strong base, such as NaH , in a non-hydroxylic solvent, such as THF. Reaction of this anion with the substituted alkyl halide (12), in which Hal is preferably I or Br, produces the oxime-ether (13).
- 25

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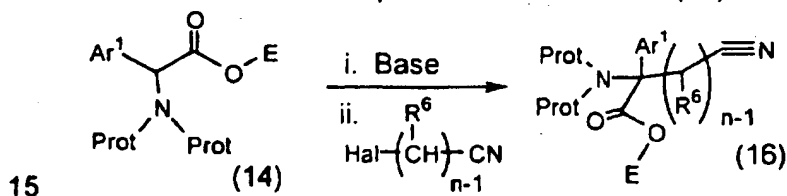
Cleavage of the oxime-ether (13) under acidic conditions, for instance, using 6N HCl at elevated temperature for 5 to 50 hours results in isolation of the ketone (2).

Further processing of (2) may be performed as described above in Method 1.

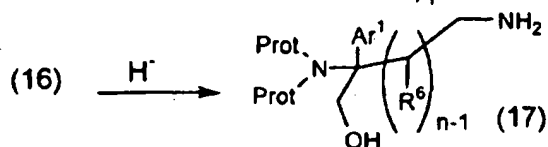
5 Method 4:

Compounds of formula I wherein X¹ is -O- and n is 2-4 can be prepared by the following method (only R⁶ is shown in the formulae, but both R⁶ and R⁷ can be present).

A diprotected aryl glycine ester, such as (14), in which "Prot" is a protecting group, preferably benzyl, and "E" is an ester group, preferably methyl or ethyl, may be converted to its anion using a strong base, such as lithium diisopropylamide, in an ether solvent, such as THF, at a temperature of about -78 °C. Reaction of the anion with a halo-nitrile (15), in which "Hal" is preferably I, at temperatures between -78 °C and 0 °C results in the protected intermediate (16).



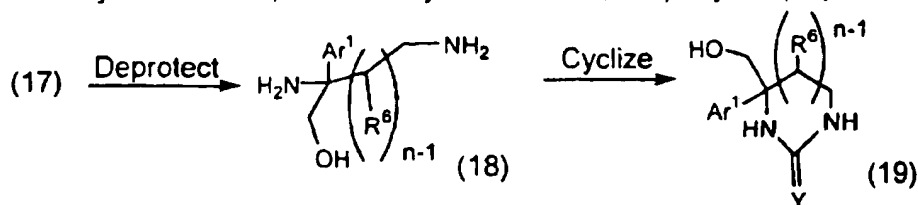
Reduction of the ester and nitrile simultaneously using a powerful hydride reducing agent, such as LAH, in an inert solvent, such as THF, at a temperature between about -78 °C and 0 °C, produces aminoalcohol (17).



Removal of the protecting groups under standard conditions (e.g. 20% Pd(OH)₂ on active carbon in methanol if "Prot" is benzyl) yields the di-amine alcohol (18) which may be cyclized using one of the reagents known to introduce a C=O, C=S

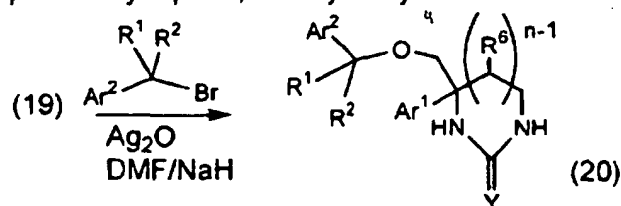
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or $-\text{SO}_2-$ group between two appropriately placed nitrogen atoms (e.g. COCl_2 ; carbonyl diimidazole; thiocarbonyl diimidazole, etc.) to yield (19).



wherein Y is $=\text{O}$ or $=\text{S}$; the thio-urea can be prepared in a similar manner.

- 5 The alcohol (19) may be converted to a compound of the invention (20), by reaction of the mono-anion prepared by using a strong base, such as NaH , with a benzyl halide in the presence of Ag_2O . "Hal" is as defined above and the solvent is preferably a polar, non-hydroxylic solvent such as DMF.

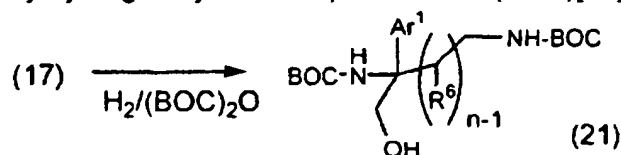


- 10 Subsequent modification by introduction of R^4 and R^5 groups may be performed as described in Method 1.

Method 4a:

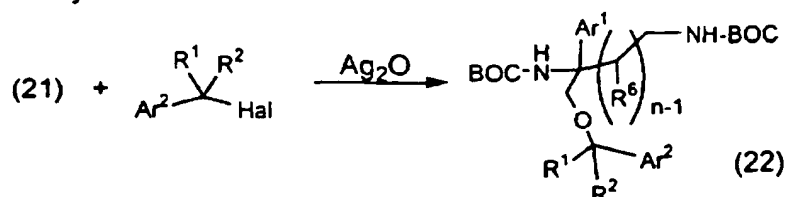
The intermediate (17), above, may also be converted to compounds of the invention via its di-BOC protected derivative (21).

- 15 Removal of the original protecting groups (if they are benzyl groups or similar) by hydrogenolysis in the presence of $(\text{BOC})_2\text{O}$ yields the di-BOC derivative (21).



Such intermediates may be converted to the ethers (22) by reaction with an aryl halide (23), preferably a bromide or iodide. Use of silver oxide (Ag_2O) as a catalyst and base is desirable.

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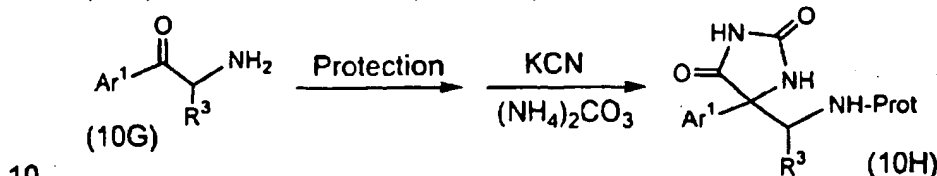


Removal of the BOC protecting groups under standard conditions (e.g. HCl /ether) produces the diamine which may be cyclized to compounds of the

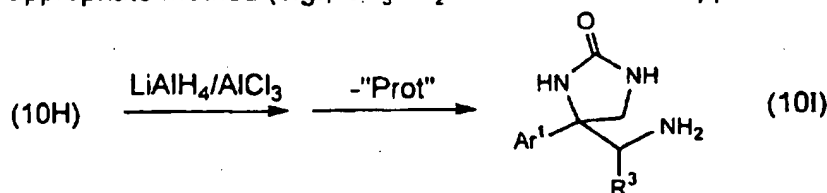
invention using the same reagents as described earlier in Method 2, e.g. COCl_2 , carbonyl diimidazole, sulfonyl diimidazole, etc.

Method 5:

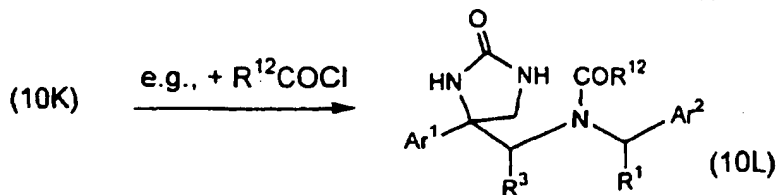
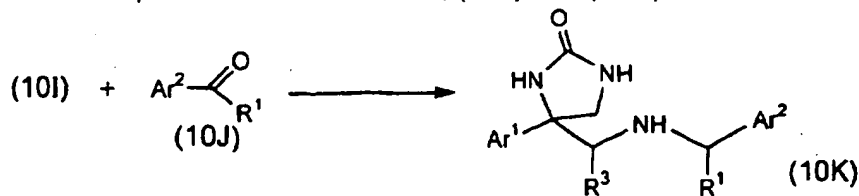
- Compounds of the invention in which X^1 is one of the nitrogen-containing groups may be synthesized from the amino ketone derivatives (10G), some of which may be commercially available while others can be synthesized by well-known literature techniques. Protection of the amino group in (10G), for instance as its BOC derivative, allows the hydantoin-forming reaction to occur to produce intermediates (10H) where "Prot" is the previously-introduced protecting group, e.g. BOC:



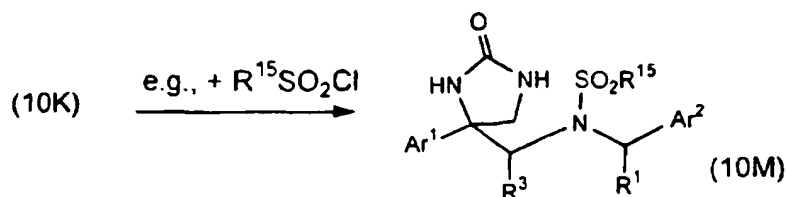
Reduction of one of the carbonyl groups preferentially, as described previously, using the $\text{LiAlH}_4/\text{AlCl}_3$ mixed reagents, followed by removal of the protecting group by an appropriate method (e.g., $\text{CF}_3\text{CO}_2\text{H}$ or HCl if it is BOC) produces the amine (10I)



- 15 Reductive alkylation of the amine group with an aldehyde or ketone (10J) under conditions described for this transformation earlier (using sodium triacetoxyborohydride) or using one of the many published procedures known to carry out this reaction (e.g. sodium borohydride in an alcohol solvent) results in the amine (10K) which may be further modified by reactions well known in the art to produce
- 20 other compounds of the invention, (10L) and (10M).



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Reactive groups not involved in the above processes can be protected during the reactions with conventional protecting groups which can be removed by standard procedures after the reaction. The following Table 1 shows some typical protecting groups:

Table 1

Group to be Protected	Group to be Protected and Protecting Group
-COOH	-COOalkyl, -COObenzyl, -COOphenyl
>NH	>NCOalkyl, >NCObenzyl, >NCOphenyl, >NCH ₂ OCH ₂ CH ₂ Si(CH ₃) ₃ , >NC(O)OC(CH ₃) ₃ , >N-benzyl, >NSi(CH ₃) ₃ ,
-NH ₂	
-OH	-OCH ₃ , -OCH ₂ OCH ₃ , -OSi(CH ₃) ₃ , or -OCH ₂ phenyl

For preparing pharmaceutical compositions from the compounds described by this invention, inert, pharmaceutically acceptable carriers can be either solid or liquid. Solid form preparations include powders, tablets, dispersible granules, capsules, cachets and suppositories. The powders and tablets may be comprised of from about 5 to about 95 percent active ingredient. Suitable solid carriers are known in the art, e.g. magnesium carbonate, magnesium stearate, talc, sugar or lactose. Tablets, powders, cachets and capsules can be used as solid dosage forms suitable for oral administration. Examples of pharmaceutically acceptable carriers and methods of manufacture for various compositions may be found in A. Gennaro (ed.), Remington's Pharmaceutical Sciences, 18th Edition, (1990), Mack Publishing Co., Easton, Pennsylvania.

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Liquid form preparations include solutions, suspensions and emulsions. As an example may be mentioned water or water-propylene glycol solutions for parenteral injection or addition of sweeteners and opacifiers for oral solutions, suspensions and emulsions. Liquid form preparations may also include solutions for intranasal administration.

Aerosol preparations suitable for inhalation may include solutions and solids in powder form, which may be in combination with a pharmaceutically acceptable carrier, such as an inert compressed gas, e.g. nitrogen.

Also included are solid form preparations which are intended to be converted, shortly before use, to liquid form preparations for either oral or parenteral administration. Such liquid forms include solutions, suspensions and emulsions.

The compounds of the invention may also be deliverable transdermally. The transdermal compositions can take the form of creams, lotions, aerosols and/or emulsions and can be included in a transdermal patch of the matrix or reservoir type as are conventional in the art for this purpose.

Preferably the compound is administered orally.

Preferably, the pharmaceutical preparation is in a unit dosage form. In such form, the preparation is subdivided into suitably sized unit doses containing appropriate quantities of the active component, e.g., an effective amount to achieve the desired purpose.

The quantity of active compound in a unit dose of preparation for treatment of respiratory diseases; inflammatory diseases; skin disorders; ophthalmological disorders; addictions; stress related disorders; obsessive/compulsive disorders; eating disorders; mania; premenstrual syndrome; central nervous system conditions; gastrointestinal disorders; bladder disorders; atherosclerosis; fibrosing disorders; obesity; Type II diabetes; pain related disorders; and genitourinary disorders; may be varied or adjusted from about 1 mg to about 1500 mg, preferably from about 50 mg to about 500 mg, more preferably from about 20 mg to about 200 mg, according to the particular application.

The actual dosage employed may be varied depending upon the requirements of the patient and the severity of the condition being treated. Determination of the proper dosage regimen for a particular situation is within the skill of the art. For convenience, the total daily dosage may be divided and administered in portions during the day as required.

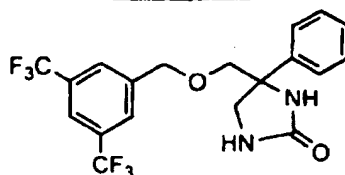
The amount and frequency of administration of the compounds of the invention and/or the pharmaceutically acceptable salts thereof will be regulated according to the judgment of the attending clinician considering such factors as age, condition and size of the patient as well as severity of the symptoms being treated. A typical

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recommended daily dosage regimen for oral administration can range from about 1 mg/day to about 1500 mg/day, in two to four divided doses.

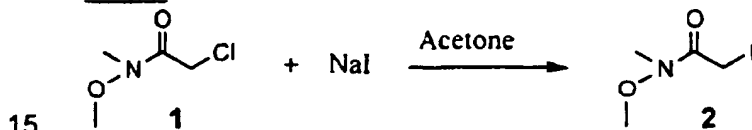
Following are examples of preparing compounds of formula I. As used herein,
 5 RT is room temperature, Me is methyl, Bu is butyl, Br is bromo, Ac is acetyl, Et is ethyl, Ph is phenyl, THF is tetrahydrofuran, EtOAc is ethyl acetate, Et₂O is ether, LAH is lithium aluminum hydride, CDI is 1,1-carbonyl diimidazole; HOBT is hydroxybenzotriazole; DEC is 1,2-diethylaminoethyl chloride; TFA is trifluoroacetic acid; Et₃N is triethylamine, MTBE is *t*-butyl methyl ether; DAST is diethylaminosulfur
 10 trifluoride.

Example 1



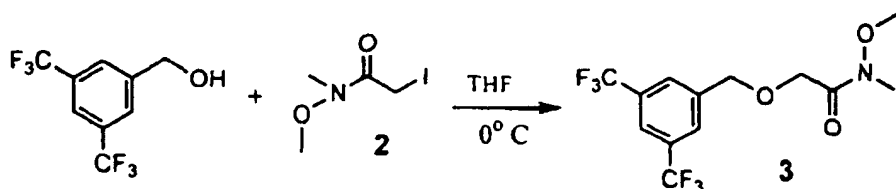
Method 1:

Step 1:



To a solution of 2-chloro-N-methylacetamide 98% (10.4 g, 74 mmol) in acetone (120 ml), NaI (12.2 g, 81.4 mmol) was added. The flask was filled with N₂ and covered with aluminum foil. After stirring at RT for 30 h, the reaction mixture was filtered. The filtrate was concentrated under vacuum to give a dark brown oil. The
 20 crude product was directly used without further purification in the next step.

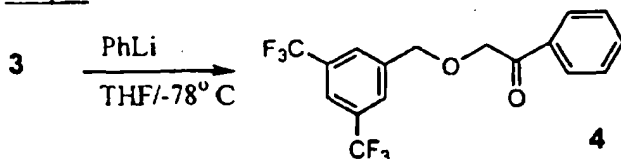
Step 2:



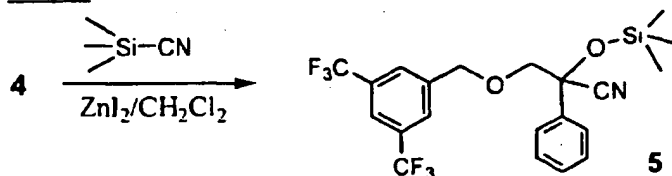
To a cooled solution of 3,5-bis(trifluoromethyl)benzyl alcohol (18.06 g, 74 mmol) in anhydrous THF (140 ml) at 0 °C, solid KN(TMS)₂ (16.24 g, 81.4
 25 mmol) was added slowly. The reaction was kept at 0 °C for 1 h. A solution of 2 in anhydrous THF (60 ml) was then added dropwise into the cooled solution. The reaction was allowed to gradually warm to RT over night under a N₂ atmosphere. After quenching with saturated NH₄Cl, the reaction was made slightly acidic with 1N HCl and extracted with EtOAc (200 ml x 4). The combined organic layer was washed

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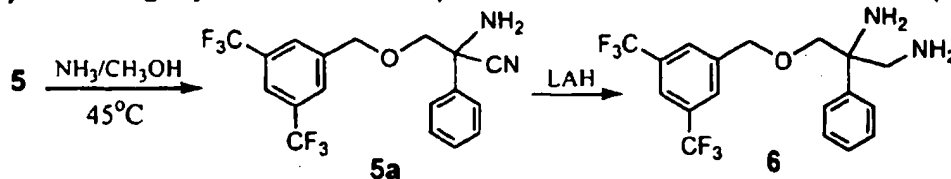
with brine (200 ml x 2), dried over MgSO_4 , filtered and concentrated. The crude material was purified by flash chromatography, eluting with 20% EtOAc in hexane to give **3** as an oil (12.3 g, 35.6 mmol, 48% yield).

Step 3:

To a cooled solution of **3** (12.2 g, 35.34 mmol) in anhydrous THF (150 ml) at -75°C , phenyl lithium (22.58 ml, 40.64 mmol) was added dropwise. The reaction mixture was kept at low temperature for 1.5 h, then the cold bath was removed and the reaction was allowed to warm up to RT under N_2 protection. The reaction was cooled in an ice-water bath and quenched with saturated NH_4Cl solution (200 ml) using a dropping funnel followed by neutralizing the aqueous solution to pH ~ 7 with 1N HCl. After stirring for 15 min, the mixture was extracted with EtOAc (200 ml x 4). The EtOAc extracts were combined, washed with brine (200 ml x 2), dried over MgSO_4 , filtered and concentrated to give a brown oil. The crude material was purified by flash grade silica gel chromatography, eluting with 15% EtOAc in hexane to give compound **4** (10.7 g, 28.44 mmol) with $\sim 80\%$ yield. FAB MS $[\text{M}+1]^+$ 263.1.

Step 4:

Trimethyl silyl cyanide (0.92 ml, 6.9 mmol) was added to a flask containing a solution of **4** (2.0 g, 5.52 mmol) in CH_2Cl_2 (11 ml), cooled with a water bath, followed by addition of ZnI_2 (88 mg, 0.276 mmol). The reaction was finished in 1 h. The insoluble ZnI_2 was filtered off and rinsed with CH_2Cl_2 . Evaporation of the solvent yielded a light yellow oil as crude product which was used in the next step.



The crude material **5** was treated with saturated $\text{NH}_3\text{-CH}_3\text{OH}$ (10 ml) and heated in an oil bath under N_2 at 45°C . After 2 h of heating, solid was filtered off and the filtrate was concentrated to give **5a**.

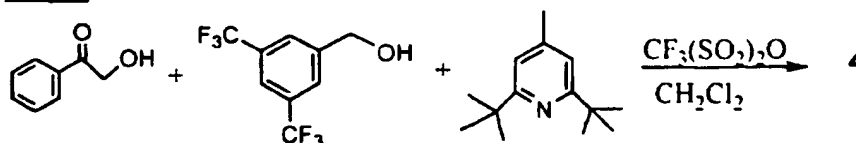
To a suspension of LAH (0.84 g, 22 mmol) in anhydrous Et_2O (40 ml) at -78°C , a suspension of **5a** in anhydrous Et_2O (40 ml) was added through an addition funnel

- 20 -

under the protection of N_2 . The reaction mixture was stirred at RT over night. After the reaction was complete, Et_2O (250 ml) was added. The reaction was quenched with saturated Na_2SO_4 solution and stirred for 2 h. The solid was filtered off and the filtrate was dried over $MgSO_4$, filtered again and treated with 4M HCl in dioxane (3 ml). Evaporation of all the volatile solvents gave the crude product, which was further purified on Biotage cartridge, eluting with 7.5% NH_4OH - CH_3OH (1:9) in 92.5% in CH_2Cl_2 to give **6** as an oil. FAB MS $[M+1]^+$ 393.1.

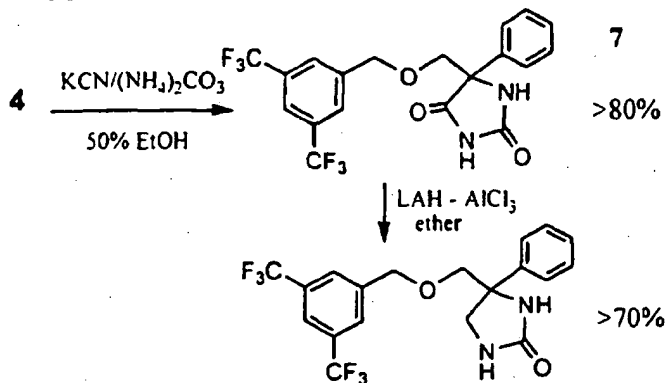
Step 5:

To a solution of **6** (0.155 g, 0.395 mmol) in anhydrous THF (12 ml) was added 3 Å molecular sieves (200 mg) and 1,1'-carbonyl diimidazole (76.6 mg, 0.474 mmol). The mixture was stirred at RT overnight under N_2 . Diluted with 200 ml EtOAc, the reaction mixture was washed with brine (100 ml x 2), dried over $MgSO_4$, filtered and concentrated. The crude material was purified by flash chromatography, eluting with 5% NH_4OH - CH_3OH (1:9) in 95% CH_2Cl_2 , to give the title compound as a solid with a 97% yield. FAB MS $[M+1]^+$ 419.1.

Method 2:**Step 1:**

A mixture of 3,5-bis(trifluoromethyl)benzyl alcohol (39.9 g, 0.163 mol) and 2,6-di-*t*-butyl-4-methyl-pyridine (68.56g, 0.33 mol) was placed in a 2 liter, 3-neck flask and vacuum dried overnight. To this green mixture was added dry CH_2Cl_2 (600 ml) (cooled with a water bath), followed by slow addition of trifluoromethylmethane sulfonic anhydride (50 g, 0.177 mol) through a dropping funnel under a N_2 atmosphere. After stirring at RT for 4 h, a solution of 2-hydroxy-acetophenone (20.36 g, 0.149 mol) in dry CH_2Cl_2 (120 ml) was added slowing through a dropping funnel under N_2 . The reaction mixture was stirred at RT for 5 days, then solid was filtered off. The filtrate was washed with brine (200 ml, 3x), dried ($MgSO_4$), filtered and concentrated to give a dark brown oil which was purified with flash grade silica gel (1 Kg), eluting with 10 % EtOAc/hexane. Compound **4** was obtained as a solid (39.63 g, 0.11 mol, 74 % yield).

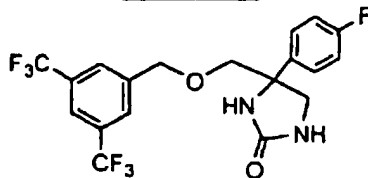
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Step 2:

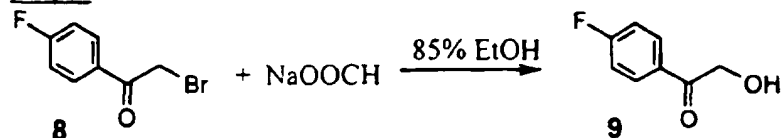
A mixture containing compound 4 (39.5 g, 109 mmol), KCN (10.64g, 163 mmol) and (NH₄)₂CO₃ (37.47 g, 380 mmol) in 50% EtOH / H₂O was heated in an oil bath at 60 °C under the N₂ atmosphere for 22 h. After cooling, ice-water (780 ml) was added and the mixture was stirred for 1 h. The white solid was filtered and rinsed with water. The solid was crystallized from hot dichloroethane. Pure compound 7 was obtained as a white solid (39g, 90 mmol, 83%).

To a one-liter round bottom flask (cooled with an ice-water bath) containing AlCl₃ (24.6g, 186 mmol) was slowly added 1 M LAH solution in Et₂O (140 ml, 140 mmol) under an N₂ atmosphere. After stirring at 0°C for 10 min, a solution of compound 7 (20g, 46.2 mmol) in dry THF (170 ml) was added slowly to the LAH-AlCl₃ mixture. It was gradually warmed to RT and stirred at RT for 3 days. After completion, the reaction mixture was diluted with THF (200 ml) and excess LAH was decomposed with saturated Na₂SO₄ and 3 N NaOH. The mixture was stirred for 1 h at RT and solid was filtered off through a celite pad. The filtrate was dried over MgSO₄, filtered and concentrated to give the title compound as a foam which was purified on flash silica gel, eluting with 3.5 % (1:9) NH₄OH/CH₃OH in 96.5 % CH₂Cl₂ to give pure title compound (16g, 40 mmol, 87% yield) as a solid.

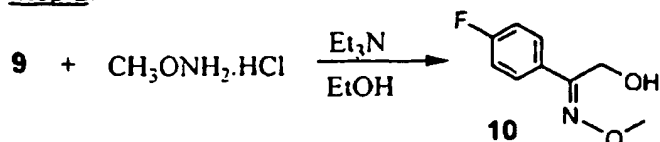
The racemic title compound was separated on a chiral column by using either a ChiralPak AS (Hexane / IPA (80:20)) or a Chiralcel OD (CH₃CN) column to give enantiomer A and enantiomer B (mp 138-140 °C). Enantiomer B: HRMS calculated for [M+1]: C₁₉H₁₇F₆N₂O₂ 419.1194, Found: 419.1190; C, H, N analysis : calculated C, 54.57; H, 3.71; N, 6.55; F 27.25 Found: C 54.55; H, 3.89; N, 6.59; F, 27.22; rotation in CH₃OH [α]_D²⁰ = -67.0 °

Example 2

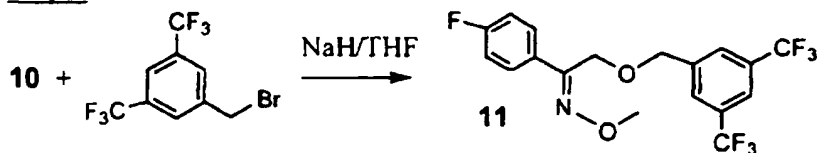
- 22 -

Part A:Step 1:

- A suspension of 2-bromo-4'-fluoroacetophenone, **8**, (30 g, 0.1368 mol) and HCOONa (59.2 g, 0.87 mol) in 85% EtOH (360 ml) was heated to 80 °C for 5h. After cooling, it was stirred at RT over night under N₂. The reaction was stopped by evaporating off the EtOH, and the residue was redissolved in brine (500 ml) and extracted with EtOAc (200 ml x 3). The combined organic layers were dried (MgSO₄), filtered and concentrated. The purified product was obtained by triturating the crude material with CH₂Cl₂, EtOAc and hexane to give **9** as a solid (18 g, 116 mmol, 85% yield).

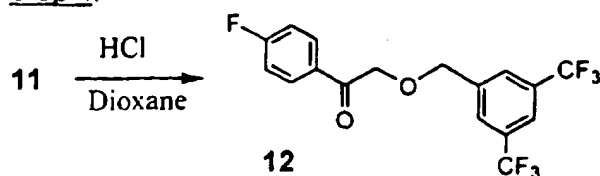
Step 2:

- To a suspension of **9** (5.0 g, 32.44 mmol) in absolute EtOH (50 ml), 3Å molecular sieves (0.8 g), CH₃NH₂.HCl (4.146 g, 48.67 mmol) and Et₃N (6.77 ml, 48.67 mmol) were added. The mixture was heated to 85 °C under N₂ for 2.5 h. The mixture was cooled to RT, the solvent was evaporated under reduced pressure, and the crude product was redissolved in EtOAc (350 ml). This organic layer was washed with brine (150 ml x 3), dried (MgSO₄), filtered and evaporated to dryness to give **10** as a solid (13.5 g, 7.36 mmol) in 97% yield.

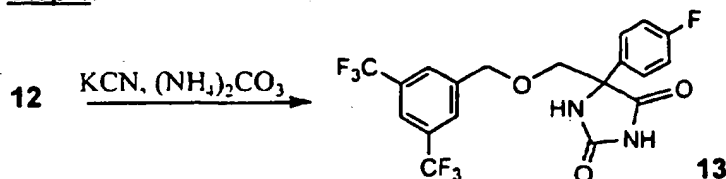
Step 3:

- To a cooled solution of **10** (12.0 g, 65.5 mmol) in anhydrous THF (60 ml) (cooled in an ice-water bath) was added 60% NaH in mineral oil (3.14g, 78.61 mmol). After stirring at RT for 15 min, the yellow suspension was treated with a solution of 3,5-bis(trifluoromethyl)benzyl bromide (13.62 ml, 72.05 mmol) in THF (13.62 ml). TLC indicated that the reaction was complete after 1.5 h at RT. Solvent was evaporated and the residue was redissolved in EtOAc (200 ml). This solution was washed with brine (50 ml x 2), dried (MgSO₄), filtered and concentrated to give **11** as an oil (26.5g, 64.77 mmol, 98% yield) FAB MS [M+1]⁺ 410.1.

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Step 4:

To a solution of 11 (26 g, 63.55 mmol) in dioxane (116 ml) was added 6N HCl (378 ml, 2.268 mol). The mixture was heated at 100 °C for 2 days. After cooling to RT, the reaction mixture was poured into ice-cold KOH solution (120 g, ~2.142 mol in 240 ml H₂O). The products were extracted with CH₂Cl₂ (200 ml x 3) from aqueous solution, dried (MgSO₄) and filtered. Solvents were evaporated to give a crude product as a brown oil. The final product was further purified by flash chromatography, eluting with 10% EtOAc in hexane to give 12 as a solid (13.41 g, 35.27 mmol, 55.5% yield).

Part B:Step 1:

To the mixture of 12 (3.23 g, 8.5 mmol) in 1:1 EtOH/H₂O (20 ml), KCN (0.83 g, 12.8 mmol) and (NH₄)₂CO₃ (2.86 g, 29.8 mmol) were added. The mixture was heated to 56 °C overnight, then cooled to RT. Iced water (60 ml) was added to the complete reaction. After 30 min, the solid was filtered off and rinsed with CH₂Cl₂ to give 13 as a white powdery product (3.12 g, 82%). FAB MS [M+1]⁺ 451.1.

Step 2:

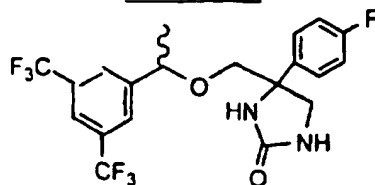
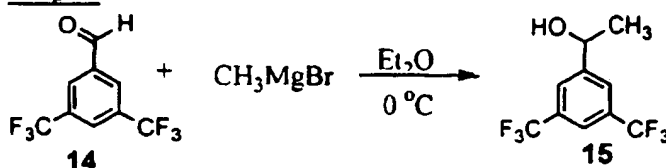
To a cooled 3-neck flask containing AlCl₃ (1.4 g, 10.7 mmol) at 0 °C, 1M LAH in Et₂O (8 ml, 8 mmol) was added dropwise via a syringe. A solution of 13 (1.2 g, 2.67 mmol) in anhydrous THF (10 ml) was added slowly and the reaction mixture was allowed to stir at RT overnight. After the reaction was complete, THF (100 ml) was added and excess LAH was decomposed with saturated Na₂SO₄ and 4N NaOH. After stirring for 1 h, the organic layer was separated, dried (MgSO₄), filtered and concentrated. The crude material was purified by flash silica gel eluting with 3.5% NH₃-CH₃OH (1:9) / 96.5% CH₂Cl₂ to give the title compound as a white foam (0.95 g, 0.22 mmol, 82% yield). FAB MS [M+1]⁺ 437.1.

The racemic compound was separated with a Chiralcel OD column, eluting with CH₃CN to give enantiomers A and B. The enantiomers A and B were reduced separately with LAH-AlCl₃ to give chiral compounds A and B. Pure chiral B was

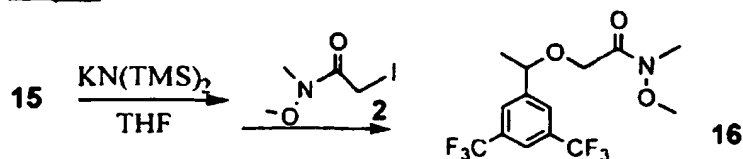
- 24 -

obtained by crystallization from hot CH₃CN and hexane. Enantiomer B: HRMS calculated for [M+1]: C₁₉H₁₆F₇N₂O₂ 437.1103, Found: 437.1100; C, H, N analysis : calculated C, 52.30; H, 3.46; N, 6.42; Found: C 52.38; H, 3.42; N, 6.31; rotation in CH₃OH [α]_D²⁰ = -55.7, mp: 130-132 °C.

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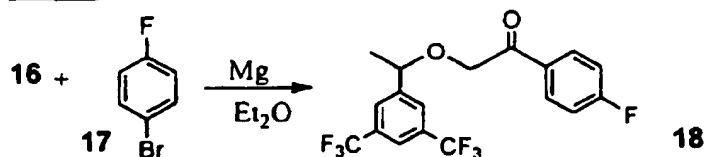
Example 3Step 1:

To the cooled solution of 3,5-bis(trifluoromethyl)benzaldehyde **14** (15 g, 62 mmol) in anhydrous Et₂O (100 ml) at 0 °C, 3 M CH₃MgBr (25 ml, 75 mmol) was added dropwise. The reaction was kept at 0 °C for another 2 h, then stirred at RT for 2 h. The reaction was quenched by pouring into ice cold saturated NH₄Cl. Routine work-up gave **15** as an off-white solid product (15.4 g).

Step 2:

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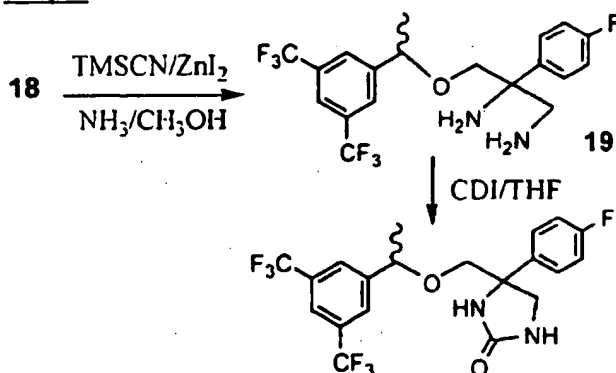
To the cooled solution of **15** (18.8 g, 72.9 mmol) in anhydrous THF (100 ml) at 0 °C was added solid KN(TMS)₂ (17.4 g, 87 mmol) portionwise. After stirring at 0 °C for 1h, the potassium salt solution of **15** was added dropwise to a solution of **2** (17.5 g, 76.4 mmol) in anhydrous THF (20 ml). The reaction was kept at 0 °C overnight and quenched with saturated NH₄Cl (300 ml). The aqueous layer was extracted with CH₂Cl₂ (300 ml x 3). The combined organic layer was washed with brine (300 ml), dried (MgSO₄), filtered and concentrated to a yellow oil (22 g). The crude product was further purified on silica gel (1:4 EtOAc/hexane) to give **16** as a pale yellow material (11 g, 31 mmol, 43% yield).

Step 3:

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- 25 -

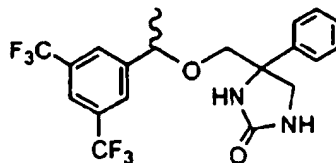
In a cooled flask (ice water bath) containing Mg turnings (1.12 g, 46.2 mmol) in anhydrous Et₂O (100 ml) was slowly added 4-bromofluoro-benzene **17** (7.7 g, 44 mmol), followed by a catalytic amount of 1,2-dibromoethane. After completion of the addition, the reaction was heated at 80 °C overnight under a N₂ atmosphere. After cooling, a solution of **16** (7.9 g, 22 mmol) in anhydrous Et₂O (40 ml) was added slowly to the Grignard solution at RT. After reaction was complete, a saturated NH₄Cl solution was slowly added under a N₂ atmosphere. The product was extracted from aqueous solution with CH₂Cl₂, combined, dried (Na₂SO₄), filtered and evaporated to give **18** as an oil. The crude material was purified on flash grade silica gel (400 g) eluting with 15% EtOAc/85% hexane to give **18** (7 g, 1.78 mmol, 87% yield). FAB MS (M+1)⁺ 395.0.

Step 4:

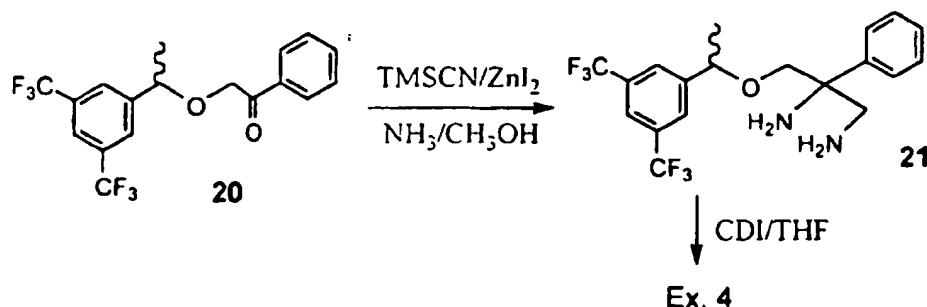
Compound **19** was prepared by an analogous method to that of Example 1 (step 4) using compound **18** in place of compound **4**.

The title compound was prepared by a method analogous to that described for Example 1 (step 5) using compound **19** in place of compound **6**. Two sets of diastereomers A and B were isolated by flash silica chromatography at compound **19** stage and these were cyclized separately to the title compounds with CDI reagent.

Diastereomer A of the title compound, FAB MS [M+1]⁺ 451.1265, calculated [M+1]⁺ 451.1257; Diastereomer B of the title compound, FAB MS [M+1]⁺ 451.1261, calculated [M+1]⁺ 451.1257.

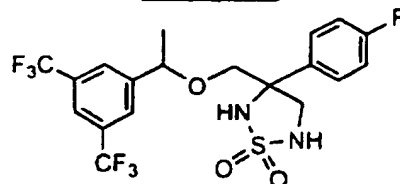
Example 4

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The title compound was prepared by an analogous method to that described for Examples 1 and 3 using α -methyl-3,5-bis(trifluoromethyl)-benzyl alcohol in place of 3,5-bis(trifluoromethyl)benzyl alcohol. Two sets of diastereomers were isolated by flash silica chromatography at compound 21 stage and cyclized separately to the title compound with CDI reagent. Diastereomer A of the title compound, FAB MS $[M+1]^+$ 433.1358, calculated $[M+1]^+$ 433.1351; diastereomer B of the title compound, FAB MS $[M+1]^+$ 433.1358, calculated $[M+1]^+$ 433.1351.

Example 5

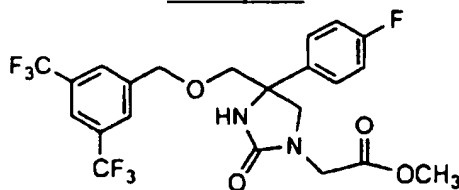


10

To a solution of compound 19 (0.22 g, 0.52 mmol) (prepared from compound 18 in Example 3) in pyridine (2 ml), sulfamide (50 mg, 0.52 mmol) was added. The mixture was refluxed for 24 h. After cooling, the reaction was diluted with EtOAc (200 ml), washed with 0.5 N HCl, brine, dried ($MgSO_4$), filtered and concentrated to give a gum which was further purified on silica gel (1:4 EtOAc/hexane) to give the title compound as an off-white solid (40 mg, 16% yield). FAB MS $[M+1]^+$ 487.3.

15

Example 6

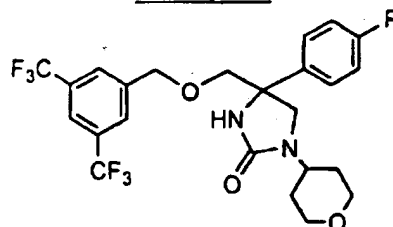
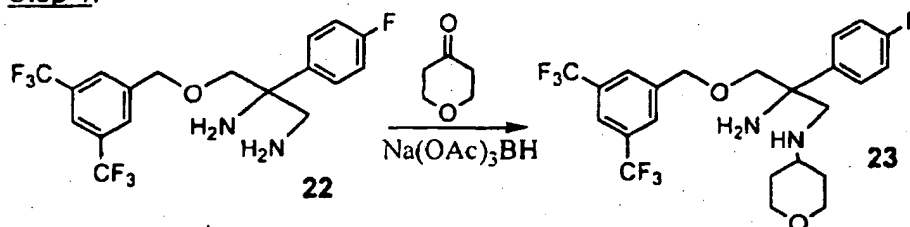


A mixture of the product of Example 2 (0.42 g, 0.96 mmol), toluene (5 ml), CH_3ONa (0.13 g, 2.3 mmol) and Bu_4NHSO_4 (3.3 mg, 0.01 mmol) was heated to 90 °C for 1h. Methyl bromoacetate (0.35 g, 2.3 mmol) was added later. After stirring at 80 °C for 48 h, the reaction was cooled to RT and product was extracted from the aqueous layer with CH_2Cl_2 (50 ml, 3x), washed with brine, and the organic layer was dried ($MgSO_4$), filtered and concentrated to give a cloudy gum (0.4 g) which was

20

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further chromatographed with 3% $\text{NH}_3\text{-CH}_3\text{OH}$ (1:9) / 97% CH_2Cl_2 to give pure title compound as a solid (35 mg, 0.07 mmol, 8% yield). Electrospray MS $[\text{M}+1]^+$ 509.1.

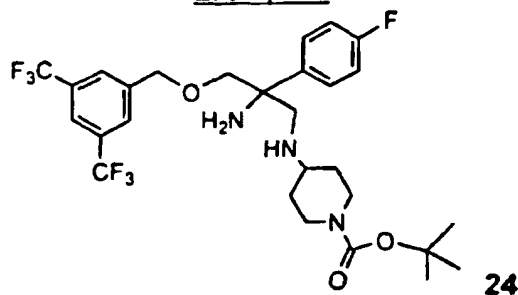
Example 75 Step 1:

To the mixture of **22** (0.205 g, 0.5 mmol) (prepared analogously to the procedures described for **6** in Example 1) in dichloroethane (4 ml) were added tetrahydro-4H-pyranone (50 mg, 0.5 mmol) and sodium triacetoxo-borohydride (0.21 mg, 1 mmol). After stirring overnight at RT, the reaction mixture was worked-up. The crude product was further purified on silica gel, eluting with 3.5% [1:9] $\text{NH}_3\text{-CH}_3\text{OH}$ in 96.5% CH_2Cl_2 . Compound **23** was obtained as an oil (0.21g, 0.43 mmol, 84% yield). FAB MS $[\text{M}+1]^+$ 495.45.

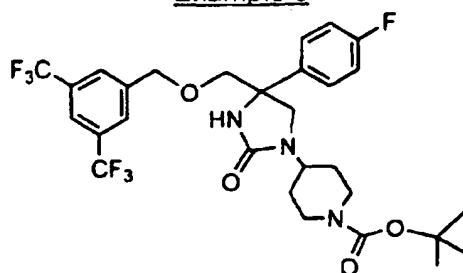
Step 2:

15 A mixture of **23** (0.15 g, 0.303 mmol) and CDI (120 mg, 0.74 mmol) in THF (5 ml) was stirred under a N_2 atmosphere at RT overnight. After work-up, a gummy material was obtained as a crude product which was then purified by chromatography, eluting with 2% $\text{NH}_3\text{-CH}_3\text{OH}$ (1:9) / 98% CH_2Cl_2 to give the title compound as a white foam (0.11 g, 0.22 mmol, 73%). FAB MS $[\text{M}+1]^+$ 521.2.

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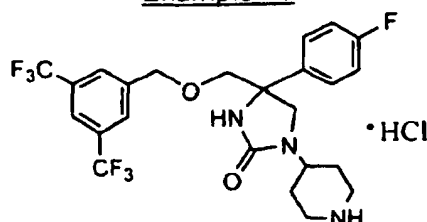
Example 8

The title compound **24** was prepared by an analogous method to that described for compound **23** in Example 7 using 1-t-butoxycarbonyl-4-piperidone in place of tetrahydro-4H-pyranone. FAB MS $[M+1]^+$ 594.1.

Example 9

The title compound was prepared from compound **24** by an analogous method to that described for the title compound of Example 7. FAB MS $[M+1]^+$ 620.2.

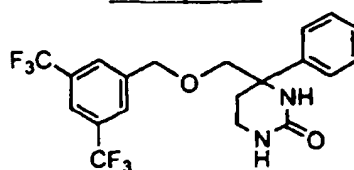
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Example 10

A solution of the product of Example 9 (95 mg, 0.153 mmol) in CH_2Cl_2 (1 ml) was treated with 4M HCl in dioxane (2 ml) for 2 h. The reaction was worked-up by evaporating solvents and excess HCl. The title compound was obtained as an off-white salt.

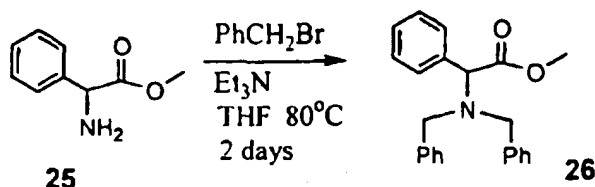
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FAB MS $[M+1]^+$ 520.3.

Example 11

Step 1:

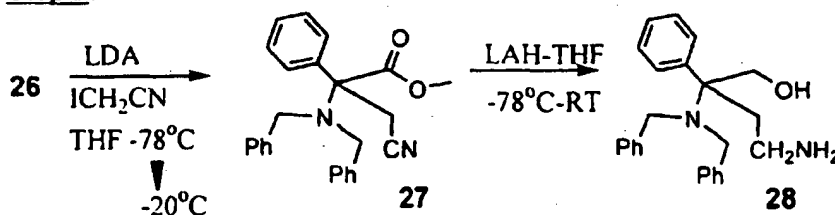
- 29 -



To a suspension of (R)-(-)-2-phenylglycine methyl ester hydrochloride salt **25** (10 g, 49.6 mmol) in anhydrous THF (30 ml) and anhydrous DMF (10 ml) at RT were added Et₃N (22.13 ml, 158.9 mmol) and benzyl bromide (14.74 ml, 123.9 mmol).

- 5 After 5 h of heating, the reaction mixture was treated with another equivalent of benzyl bromide (5.89 ml, 49.6 mmol) and Et₃N (6.9 ml, 49.6 mmol). The solution was heated at 80 °C under a N₂ atmosphere overnight. Additional benzyl bromide (6.9 ml, 49.6 mmol) was added next day and heating continued at 80 °C overnight to complete the reaction. After completion, the reaction mixture was cooled to RT, and
- 10 poured into a separatory funnel containing NaHCO₃ solution and EtOAc. The aqueous layer was extracted with EtOAc (200 ml x 3). The combined organic layer was washed with brine (300 ml), dried over MgSO₄, filtered and concentrated. Crude product was purified by chromatography (3% EtOAc in hexane) to give **26** as an oil (.836 g, 22.62 mmol, 46 % of yield). Electrospray MS [M+1]⁺ 346.1.

15 Step 2:

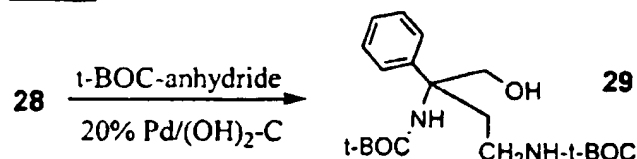


- To a cooled solution of **26** (18 g, 52.11 mmol) in anhydrous THF (125 ml) at -78 °C, a 2M LDA (32.6 ml, 65.2 mmol) solution in THF/n-heptane was added slowly. The reaction mixture was kept at this low temperature for another 2h under a N₂
- 20 atmosphere, then treated with 95% ICH₂CN (4.97 ml, 65.2 mmol) dropwise through a syringe. The reaction was stirred at -78 °C for 4h and -20 °C overnight. A mixture of chilled saturated aqueous NH₄Cl solution was added to quench the reaction. The separated aqueous layer was extracted with EtOAc (200 ml x 3). The combined organic layers were washed with brine (300 ml), dried over MgSO₄, filtered and
- 25 concentrated. The residue was purified by flash chromatography (15% EtOAc in hexane) to afford **27** as an oil (10.24 g, 51.2%). Electrospray MS [M+1]⁺ 385.1.

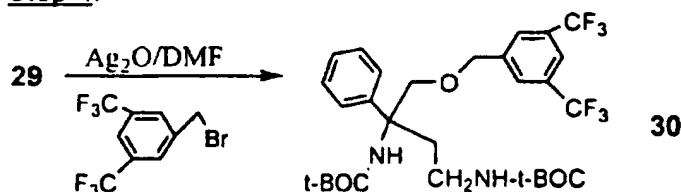
- To a solution of **27** (6.0 g, 15.6 mmol) in anhydrous THF (100 ml) was added 1 M LAH in THF (100ml, 100 mmol) slowly at -78 °C under a N₂ atmosphere. The reaction mixture was let to stir from -78 °C to RT overnight. After completion, the
- 30 reaction was diluted with THF (100 ml) and quenched (cooled in an ice-bath) with a

- 30 -

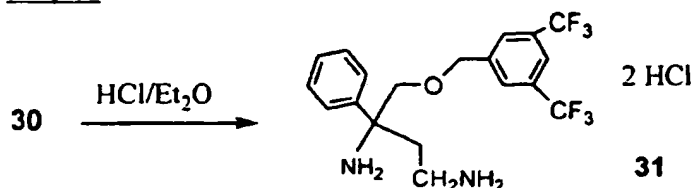
Na₂SO₄ saturated aqueous solution slowly through a dropping funnel. The mixture was stirred at RT for 1h, then filtered and concentrated. The crude product was triturated with CH₃OH and EtOAc and filtered. The solid (2.24 g) was the desired pure compound. The filtrate was further purified by flash grade silica gel, eluting with
 5 6% NH₄OH-CH₃OH (1:9) / 94 % CH₂Cl₂ to give compound **28** (1.3 g) as a solid with a total yield of 64 %.

Step 3:

To a solution of **28** (1.27 g, 3.523 mmol) in CH₃OH (150 ml) was added t-BOC anhydride (1.8 g, 8.24 mmol) and 20% Pd(OH)₂ on carbon (0.254 g). The mixture
 10 was hydrogenolyzed at 50 psi in a Parr Shaker overnight. After completion, excess catalyst was filtered and rinsed with CH₃OH. Solvent was concentrated and gave the crude product **29**, which was further purified on flash grade of silica gel, eluting with 5% NH₃-CH₃OH (1:9) in 95% CH₂Cl₂ to give **29** as a white solid (1.14 g, 2.99 mmol)
 15 with a 85 % of yield. Electrospray MS [M+1]⁺ 381.1.

Step 4:

To a solution of **29** (2.4 g, 6.306 mmol) (from another batch) in anhydrous DMF (20 ml) was added 4 Å molecular sieves (1 g), 3,5-bis(trifluoromethyl)benzyl bromide (1.736 ml, 9.459 mmol) and Ag₂O (2.998 g, 12.61 mmol) under a N₂ atmosphere.
 20 The reaction was stirred at RT in the dark overnight. EtOAc (300 ml) was added, and the mixture was washed with brine (100 ml x 2), dried over MgSO₄, filtered and evaporated to give **30** as a yellow oil. It was purified by chromatography, eluting with 0.75% NH₃-CH₃OH (1:9) / 99.5 % CH₂Cl₂ to give **30** as a solid (2.4 g, 3.95 mmol, 63%
 25 yield). Electrospray MS [M+1]⁺ 607.1.

Step 5:

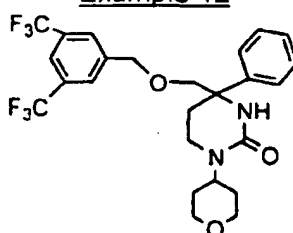
- 31 -

To a solution of **30** (1.38 g, 2.27 mmol) in CH_2Cl_2 (8.5 ml) was added 2N HCl / Et_2O (8.5 ml, 17 mmol). The solution was stirred at RT overnight under a N_2 atmosphere. After the reaction was complete, all solvents were evaporated to give compound **31** as a solid. Crude yield: 1.08 g, 99.6%.

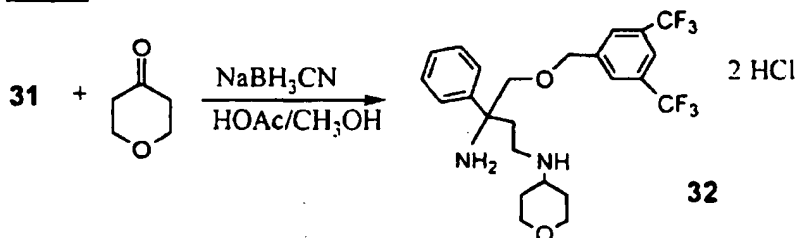
5 Step 6:

- To a suspension of **31** (1.07 g, 2.233 mmol) in anhydrous THF (30 ml) was added 4 Å molecular sieves (1.0 g), Et_3N (0.653 ml, 4.69 mmol) and CDI (0.434 g, 2.679 mmol) at 0 °C. The reaction mixture was stirred at ambient temperature overnight under a N_2 atmosphere. After completion, the reaction mixture was
- 10 evaporated and the residue was partitioned between EtOAc and water (300 ml). The organic layer was combined and washed with brine (50 ml x 3), dried over MgSO_4 , filtered and concentrated to give a clear oil of the title compound. The crude product was further purified on flash grade silica gel, eluting with 5% NH_4OH -MeOH (1:9) / 95 % CH_2Cl_2 to give the title compound as a solid (0.46 g, 1.06 mmol) in a 48% of yield.
- 15 Electrospray MS $[\text{M}+1]^+$ 433.1.

Example 12



Step 1:



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- To a cooled solution of **31** (0.5 g, 1.044 mmol) in CH_3OH (14.65 ml) was added Et_3N (0.29 ml, 2.088 mmol), Na_2SO_4 (200 mg) and tetrahydro-4H-pyran-4-one (98 μl , 1.044 mmol) at 0 °C under N_2 atmosphere. After
- 25 1 h, NaBH_3CN (103.5 mg, 1.566 mmol) and HOAc (125.6 μl , 2.088 mmol) were added. The reaction was stirred at 0 °C for 3 h and 1N NaOH was added until solution reached pH=10. The volatile solvent was evaporated *in vacuo*. The residue was redissolved in CH_2Cl_2 (300 ml) and washed with aqueous mixture of saturated NaHCO_3 - NaCl - H_2O . The combined organic layer was dried over MgSO_4 , filtered and evaporated to give **32** as an oil. The crude product was passed through 50 g of flash

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grade silica gel, eluting with 5% $\text{NH}_3\text{-CH}_3\text{OH}$ (1:9) / 95 % CH_2Cl_2 to give **32** as an oil (0.28 g, 0.57 mmol, 55%). Part of this material (100 mg) was redissolved in CH_2Cl_2 and two equivalents of HCl-ether solution was added. After stirring for 10 min, solvents were evaporated to give HCl salt of compound **32** as a solid. Electrospray

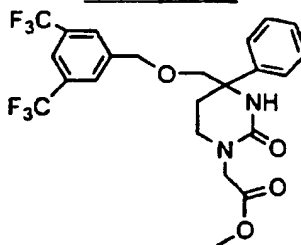
5 MS $[\text{M}+1]^+$ 491.1.

Step 2:

To a solution of **32** (180 mg, 0.367 mmol) in anhydrous THF (7.0 ml) was added 4Å molecular sieves (300 mg) and CDI (71.4 mg, 0.44 mmol). The reaction mixture was stirred at RT for over 50 h. A small amount of NaHCO_3 solution was
10 added to quench the reaction. The cloudy mixture was diluted with EtOAc (200 ml). The organic layer was washed with brine (50 ml x 3), dried over MgSO_4 , filtered and concentrated. The crude product was purified on 30 g of flash grade silica gel, eluting with 5% $\text{NH}_3\text{-CH}_3\text{OH}$ (1:9) / 95 % CH_2Cl_2 to give the title compound as a solid (0.118 g, 0.23 mmol, 63%). FAB MS $[\text{M}+1]^+$ 517.1.

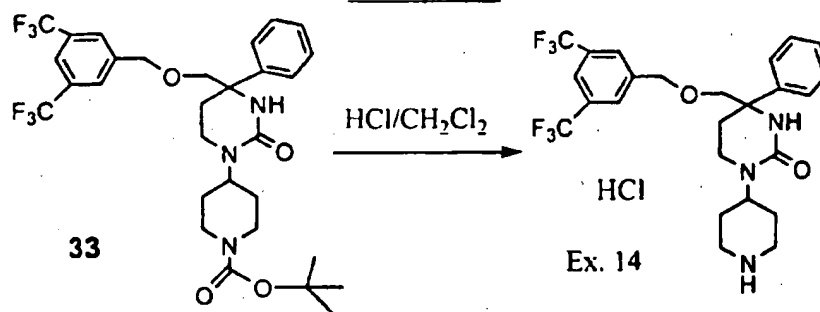
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Example 13



To a cooled solution of the product of Example 11 (0.36 g, 0.832 mmol) in anhydrous DMF (2 ml) at 0 °C was added 60% NaH (40 mg, 0.998 mmol) in mineral oil. After stirring 30 min, methylbromo acetate (88 μl , 0.915 mmol) was added. The
20 reaction was stirred at 0 °C to RT overnight under a N_2 atmosphere. EtOAc (300 ml) was added and the mixture was washed with brine (100 ml x 3), dried over MgSO_4 , filtered and concentrated to give a crude material. It was separated on flash grade silica gel (50 g), eluting with 5% $\text{NH}_3\text{-CH}_3\text{OH}$ (1:9) in 95% CH_2Cl_2 to give the title compound as a gummy oil (15 mg, 0.029 mmol, 3.5%). Electrospray MS $[\text{M}+1]^+$
25 505.1.

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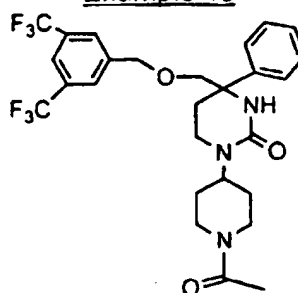
Example 14

Compound **33** was prepared by an analogous method to that described in Examples 8 and 9 using compound **31** as starting material in place of compound **22**.

5 **Electrospray MS $[\text{M}+1]^+$ 616.1.**

To a solution of **33** (328 mg, 0.533 mmol) in CH_2Cl_2 (2.66 ml) was added 2N $\text{HCl-Et}_2\text{O}$ (2.66 ml, 5.33 mmol). The reaction mixture was stirred at RT for 18h under a N_2 atmosphere. Crude title compound was obtained by evaporating off the solvents and used directly in the following step. FAB MS $[\text{M}+1]^+$ 516.1.

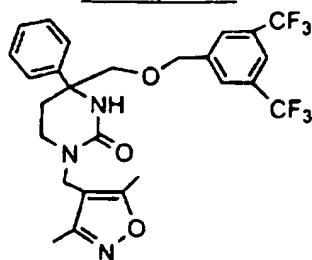
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Example 15

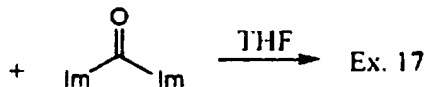
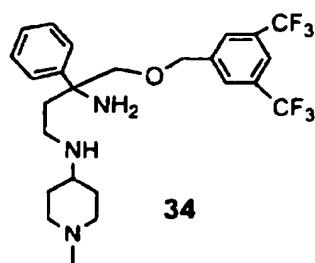
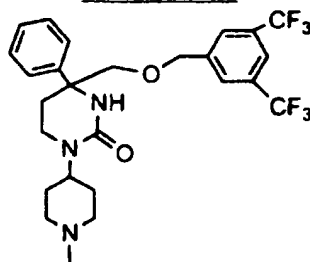
To a solution of the product of Example 14 (110 mg, 0.199 mmol) in anhydrous CH_2Cl_2 (2.2 ml) was added Et_3N (61 μl , 0.438 mmol), followed by the addition of HOAc (12 μl , 0.2 mmol), HOBT (27 mg, 0.2 mmol) and DEC (46 mg, 0.24 mmol).

- 15 The reaction mixture was stirred at RT for 3 h under a N_2 atmosphere. After completion, CH_2Cl_2 (200 ml) was added and the reaction mixture was washed with brine (50 ml x 2), dried over MgSO_4 , filtered and evaporated to give the title compound as a foam. The crude material was purified on flash grade silica gel, eluting with 5 % $\text{NH}_4\text{OH} / \text{CH}_3\text{OH}$ (1:9) / 95 % CH_2Cl_2 to afford the title compound as a
- 20 white solid (90 mg, 0.16 mmol, 81%). Electrospray MS $[\text{M}+1]^+$ 558.1.

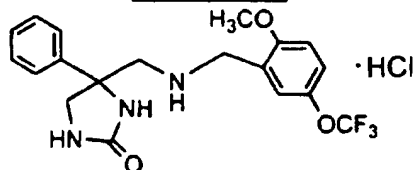
- 34 -

Example 16

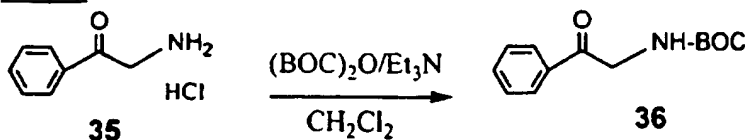
The title compound was prepared by an analogous method to that described in Example 13 using 4-chloromethyl-3, 5-dimethyl-isoxazole in place of methylbromo acetate and 60% NaH in mineral oil (1.5 equivalent). Electrospray MS $[M+1]^+$ 542.1.

Example 17

The title compound and compound 34 were prepared by a method analogous to that described in Examples 12 using 1-methyl-4-piperidone in place of tetrahydro-4H-pyran-4-one. Electrospray MS $[M+1]^+$ 530.1.

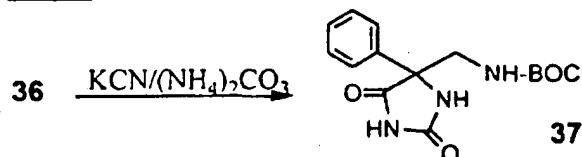
Example 18

15 Step 1:

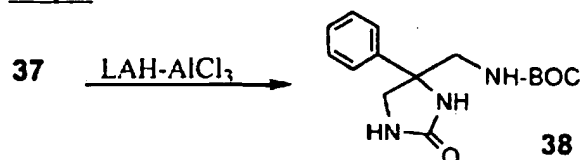


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To a solution of **35** (14.8 g, 86.23 mmol) and Et₃N (36 ml, 258.69 mmol) in CH₂Cl₂ (200 ml), (BOC)₂O (20.70 g, 94.85 mmol) was added at RT. Stirred for 18 h, the reaction was stopped by diluting with CH₂Cl₂ (100 ml) and transferred into a separatory funnel. The organic mixture was washed with water (100 ml x 2), dried
5 over Na₂SO₄, filtered, and concentrated to give a light yellow solid **36** (20 g, 85 mmol, 99%). This crude product was used without purification in the following step.

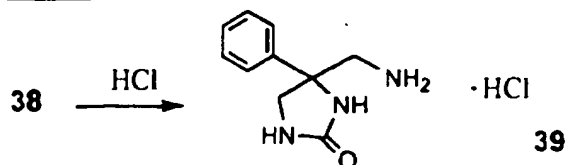
Step 2:

In a flask containing **36** (20.28 g, 86.195 mmol), (NH₄)₂CO₃ (28.99 g, 301.68 mmol) and KCN (8.42 g, 129.29 mmol) was added a (1:1) mixture of EtOH and water (180 ml). The mixture was heated to 56 °C under N₂ atmosphere for 48 h. The reaction was cooled with an ice bath and then the precipitate was filtered off. The residue was rinsed with water, then hexane, and air dried to give **37** as a white solid (20 g). The filtrate was diluted with EtOAc (500 ml) and the organic layer was
15 separated, dried over Na₂SO₄, filtered and concentrated. The solid was triturated with Et₂O then CH₂Cl₂, and dried in air to give additional **37** (4.8 g) as a solid (total 24.8 g, 81.2 mmol, 94%).

Step 3:

To a 3-neck flask equipped with a N₂ inlet was added AlCl₃ (10.48 g, 78.60 mmol). An ice-bath was placed under the flask. A solution of 1M LAH in Et₂O (59 ml, 59 mmol) was added dropwise to the cooled flask. After 10 min, a solution of hydantoin **37** (6 g, 19.65 mmol) in anhydrous THF (100 ml) was added to the reaction mixture slowly. The reaction was kept at 0 °C for additional 15 min then gradually
25 warmed up to RT overnight. In a cooling bath, the reaction was carefully quenched with water (3 ml), followed by 15% NaOH (3 ml) and water (9 ml) under an atmosphere of N₂. Stirred for 15 min, the reaction mixture was filtered and rinsed with EtOAc, the THF. The filtrate was transferred to a separatory funnel containing water (50 ml). The organic layer was separated, dried over Na₂SO₄, filtered, and
30 concentrated to give **38** as a white solid (4.5 g, 15.44 mmol, 79%). The crude product was used without further purification. FAB MS [M+1]⁺ 292.1.

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Step 4:

- To a solution of **38** (0.9 g, 3.09 mmol) in CH_3OH (20 ml), a 4 N HCl (8 ml, 31 mmol) solution in dioxane was added. After stirring at RT for 2 h, the reaction was stopped by evaporating off the volatile solvents. CH_3OH was added twice with subsequent evaporation. A white foam (0.65 g, 2.85 mmol, 92 % yield) was obtained as crude product, which was used directly in the following step.

Step 5:

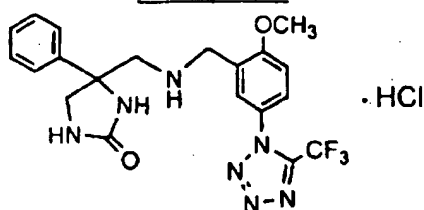
- To a solution of alcohol **40** (0.5 g, 2.43 mmol) in dioxane (2.5 ml), water (0.15 ml) was added, followed by Cs_2CO_3 (3.96 g, 12.15 mmol) and CH_3I (0.45 ml, 7.29 mmol). The mixture was heated to 72 °C in a sealed tube for 3 h. The reaction mixture was then cooled to RT and concentrated. The residue was flashed through silica gel, eluting with 40 % EtOAc/ hexane. Compound **41** was obtained as an oil (0.52 g, 2.36 mmol, 97 % yield).

Step 6:

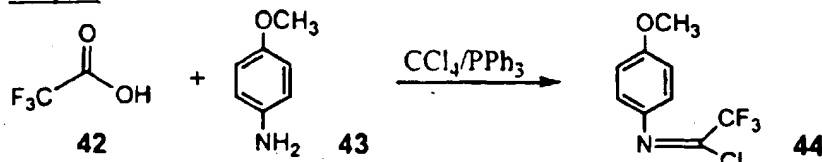
- To a solution of **39** (0.28 g, 1.23 mmol) in (9:1) of trifluoroethanol / Et_3N (10 ml), aldehyde **41** (0.27 g, 1.23 mmol) was added, followed by 3 Å molecular sieves (1.5 g) and sodium triacetoxyborohydride (0.91 g, 4.31 mmol). This turbid reaction was made clear by adding dichloroethane (15 ml). After 18 h, the reaction mixture was filtered and rinsed with EtOAc (200 ml) and CH_3OH (200 ml). The combined filtrate was concentrated. The residue was redissolved in EtOAc (250 ml) and washed with saturated NaHCO_3 (100 ml). The organic layer was dried over Na_2SO_4 , filtered and concentrated. The crude material was purified on flash silica gel, eluting with EtOAc / Et_3N (9:1) to give the title compound as an oil (0.3 g, 0.76 mmol, 62 % yield), FAB MS $[\text{M}+1]^+$ 396.1. The compound was treated with one equivalent of 1 N HCl- Et_2O solution and isolated as a HCl salt.

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Example 19

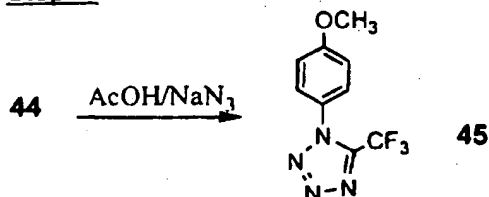


Step 1:



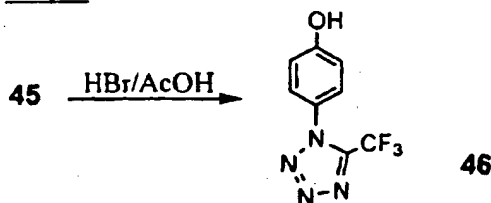
- 5 To a solution of triphenyl phosphine (99.5 g, 379.43 mmol) in CCl_4 (51 ml), Et_3N (16.98 ml, 121.80 mmol) and trifluoro acetic acid, **42**, (7.8 ml, 101.08 mmol) were added. After stirring at 0°C for 10 min, a solution of
- 10 *p*-anisidine **43** (15 g, 121.80 mmol) in CCl_4 (51 ml) was added. The reaction mixture was refluxed for 3 h. Cooled down to RT, the reaction mixture was filtered and rinsed with hexane until there was no yellow color filtered off. The filtrate was concentrated and the brown residue was distilled at 110°C to give the desired product **44** (17 g, 71.55 mmol, 59 % yield) as a yellow oil.

Step 2:



- 15 To a solution of compound **44** (7 g, 29.46 mmol) in AcOH (110 ml) was added NaN_3 (6.13 g, 94.27 mmol). The reaction mixture was heated at 70°C overnight. After cooling to RT, the reaction mixture was filtered and the filtrate was diluted with CH_2Cl_2 (200 ml) and washed with saturated NaHCO_3 (100 ml) solution, water (100 ml) and dried over Na_2SO_4 , filtered and concentrated. Compound **45** (7 g, 28.67 mmol, 98
- 20 % yield) was obtained and used directly without further purification.

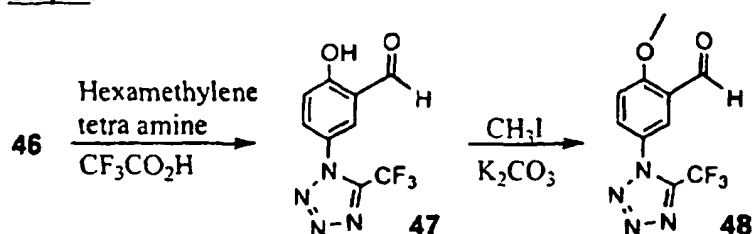
Step 3:



To the flask containing compound **45** (15 g, 61.43 mmol) and AcOH (100 ml) was added 48% aqueous HBr (100 ml). The reaction mixture was heated to 100°C

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for 48 h. After cooling to RT, the reaction mixture was concentrated to half of its volume on the rotary evaporator and extracted with EtOAc (200 ml x 2). The combined organic layer was washed with water (150 ml x 3), dried over Na₂SO₄, filtered and concentrated to give **46** (8.0 g, 34.76 mmol, 57 % yield) which was used without further purification.

Step 4:

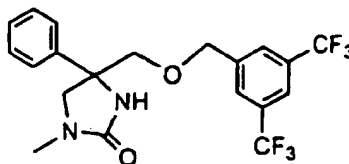
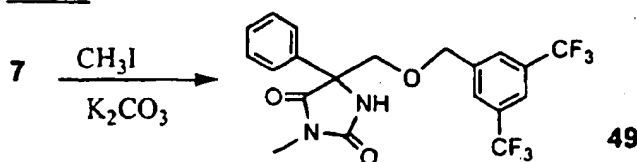
To the solution of phenol **46** (4 g, 17.39 mmol) in TFA (40 ml) was added hexamethylene tetramine (11.2 g, 79.89 mmol). The reaction mixture was heated at 70 °C under a N₂ atmosphere for 48 h. Cooled to RT, the reaction mixture was poured into 2 N H₂SO₄ (100 ml). This mixture was then extracted with EtOAc (200 ml, 2x), dried over Na₂SO₄, filtered and concentrated to give compound **47** (3.4 g, 13.17 mmol, 76 % yield) as a crude product which was used in the next step directly.

Compound **47** (3.4 g, 13.17 mmol) was dissolved in anhydrous DMF (20 ml). To this solution was added K₂CO₃ (3.6 g, 26.34 mmol) and CH₃I (1.7 ml, 26.34 mmol). The reaction mixture was stirred at RT for 3 h and then poured into a separatory funnel containing water (125 ml) and EtOAc (250 ml). The organic layer was washed with water (100 ml x 2), brine (100 ml) and concentrated. The crude product was purified by flash silica gel, eluting with hexane/EtOAc (9:1) to give **48** as a white solid (3 g, 1.02 mmol, 84%).

Step 5:

To a solution of hydrochloride salt of **39** (0.25 g, 1.098 mmol), a (9:1) mixture of trifluoroethanol/Et₃N (20 ml) and aldehyde **48** (0.3 g, 1.098 mmol) was added followed by 3Å molecular sieves (1.5 g). The reaction mixture was stirred under N₂ protection for 0.5 h before sodium triacetoxy borohydride (0.81 g, 3.83 mmol) was added. After 18 h of stirring, the reaction was filtered and rinsed with a (1:1) mixture of CH₃OH/EtOAc (200 ml). The filtrate was concentrated and purified with flash grade silica gel, eluting with EtOAc/Et₃N (9:1) to give the title compound (0.3g, 0.67 mmol, 61% yield) FAB MS [M+1]⁺ 448.1. The compound was treated with one equivalent of 1N HCl-Et₂O solution and isolated as a HCl salt.

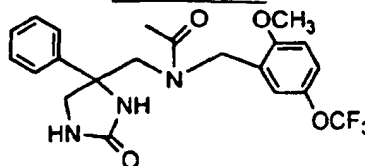
- 39 -

Example 20Step 1:

- 5 To the solution of hydantoin compound **7** (0.1 g, 0.23 mmol) (Example 1, method 2) in anhydrous DMF (1 ml), K_2CO_3 (45 mg, 0.322 mmol) was added. After stirring at RT for 25 min, CH_3I (29 μ l, 0.46 mmol) was added. After stirring at RT overnight, the reaction mixture was quenched by pouring into EtOAc (100ml). The organic layer was washed with a saturated K_2CO_3 (100 ml x 2) solution, dried over
- 10 Na_2SO_4 , filtered and concentrated. The crude product was purified by flash chromatography (EtOAc/hexane 1:4) to give compound **49** as a solid (67 mg, 0.17 mmol, 74 % yield). FAB MS $[M+1]^+$ 447.1.

Step 2:

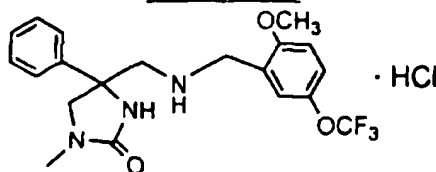
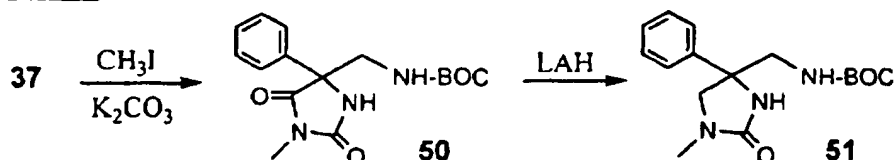
- 15 To a 2-neck flask containing $AlCl_3$ (0.37 g, 2.77 mmol), 1M LAH (2.07 ml, 2.07 mmol) in Et_2O was added slowly. Stirred for a few minutes, a solution of hydantoin **49** (0.31 g, 0.69 mmol) in anhydrous Et_2O (5 ml) was added to the white suspension. The reaction mixture was left at RT for
- 48 h and quenched by adding water (1 ml), dropwise, followed by 15% NaOH (1 ml). The white precipitate was filtered and rinsed with EtOAc. The filtrate was washed
- 20 with brine (50 ml), dried over Na_2SO_4 , filtered and concentrated. The crude product was purified by flash chromatography (2% CH_3OH in CH_2Cl_2) and the title compound was isolated as a solid (0.22 g, 0.51 mmol, 74 % yield). FAB MS $[M+1]^+$ 433.1.

Example 21

- 25 To the mixture of the compound of Example 18 (0.05 g, 0.126 mmol) in CH_2Cl_2 (2 ml), Et_3N (35 μ l, 0.252 mmol) and CH_3COCl (18 μ l, 0.252 mmol) were added. The reaction mixture was left at RT overnight and flashed through silica gel, eluting with EtOAc/ Et_3N (9:1), to give the title compound as a foam (35 mg, 0.075 mmol, 60 %

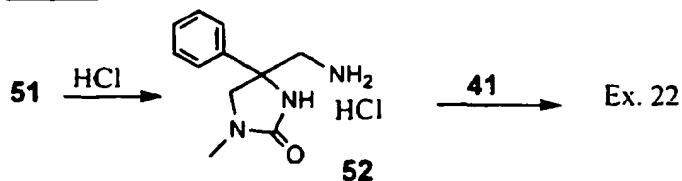
- 40 -

yield).

FAB MS $[M+1]^+$ 438.1.Example 225 Step 1:

To the solution of **37** (2 g, 6.55 mmol) in anhydrous DMF (12 ml), K_2CO_3 (1.27 g, 9.19 mmol) was added. After stirring at RT for 0.5 h, CH_3I (0.82 ml, 13.17 mmol) was added. After stirring at RT overnight, the reaction was diluted with EtOAc (150 ml) and washed with water (100 ml x 2), dried over Na_2SO_4 , filtered, and concentrated to give a white solid as the crude **50**. The crude material was further purified on the flash grade silica gel, eluting with 20 % EtOAc in hexane to give pure **50** as a solid (1.88g, 5.89 mmol, 90 % yield).

To a cooled flask containing $AlCl_3$ (3.14 g, 23.55 mmol) at 0 °C, 1M solution of LAH in anhydrous THF (18 ml 18 mmol) was added slowly. The resulting white slurry was treated with a solution of hydantoin **50** (1.88 g, 5.89 mmol) in anhydrous THF (40 ml). The reaction mixture was stirred at RT for 3 days. Water (2 ml) was added carefully to quench the remaining reactants, followed by 15% NaOH (2 ml) and water (6 ml). The mixture was then poured into a separatory funnel containing EtOAc (200 ml), and the organic layer was washed with water (100 ml x 2), dried over Na_2SO_4 , filtered and concentrated. The crude product **51** (1.28 g, 4.19 mmol, 72 % yield) was used directly in the next step.

Step 2:

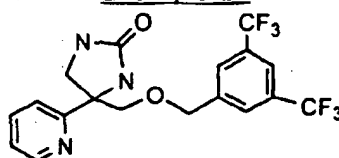
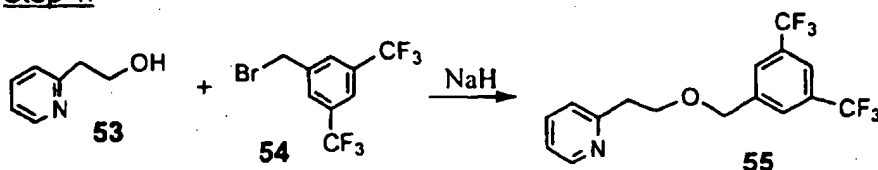
The mixture of BOC compound **51** (1.28 g, 4.19 mmol) in 4 M HCl solution of dioxane (25 ml, 100 mmol) was stirred at RT for 18 h. After completion, all solvents were evaporated off and the residue was dried by azeotropic distillation with CH_3OH twice. The foamy crude product **52** (0.1 g, 0.487 mmol) was dissolved in a solution of CF_3CH_2OH/Et_3N 9:1 (10 ml). To this clear solution, aldehyde **41** (0.107g, 0.487

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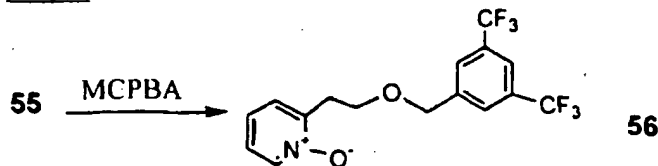
mmol) was added, followed by 3Å molecular sieves (0.7 g). After stirring at RT for 45 min, NaBH(OAc)₃ (0.36g, 1.7 mmol) was added. The reaction was stirred overnight at RT. After reaction was complete, the mixture was filtered and washed with CH₃OH and EtOAc. Evaporation of all solvents gave the title compound as a crude product which was further purified by silica gel chromatography with EtOAc/hexane (1:1) as eluting solvents to give the title compound (80 mg, 0.195 mmol, 40 %) as an oil. FAB MS [M+1]⁺ 410.1.

The compound was treated with one equivalent of 1 N HCl – Et₂O solution and isolated as the HCl salt.

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Example 23Step 1:

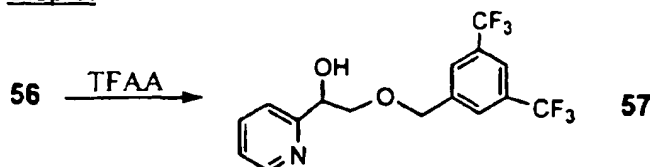
A solution of the alcohol **53** (10 g, 81.2 mmol) in anhydrous DMF (50 ml) was cooled to 0 °C and treated with 60% NaH (3.6 g, 89.3 mmol) in mineral oil portionwise. The cooling bath was removed after 30 min and the reaction was stirred at RT for 2 h. The reaction mixture was cooled again to 0 °C and then treated with bromide **54**. After stirring for 4 h, the reaction was quenched by pouring into EtOAc (250 ml). The organic mixture was washed with water (200 ml, x 4), brine (50 ml), dried over Na₂SO₄, filtered and concentrated. The crude product was purified on a Biotage 75S column, eluting with hexane/EtOAc (5:1) to give **55** (22.2 g, 63.6 mmol, 78 % yield). Electrospray MS [M+1]⁺ 350.1.

Step 2:

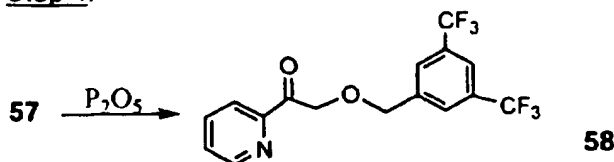
A solution of **55** (10 g, 28.6 mmol) in CH₂Cl₂ (50 ml) was treated with MCPBA (6.2 g, 28.6 mmol). After 2 h, additional 2 g of MCPBA was added and the mixture was stirred for 6 h. The mixture was poured into EtOAc (250 ml) and washed with saturated aqueous Na₂S₂O₃ (200 ml), saturated aqueous NaHCO₃ (200 ml), and

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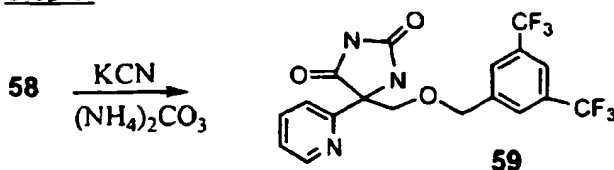
brine, dried over Na_2SO_4 , filtered and concentrated to give the N-oxide **56** as a lightly colored solid (10.18 g, 27.9 mmol, 97.5% yield). Electrospray MS $[\text{M}+1]^+$ 366.1.

Step 3:

- 5 A mixture of N-oxide **56** (6.0 g, 16.43 mmol) and trifluoroacetic anhydride (17.3 g, 82.2 mmol) in CH_2Cl_2 (75 ml) was heated under reflux for 4 h and then stirred at RT overnight. The reaction mixture was concentrated *in vacuo*. The residue was dissolved in THF (40 ml) and saturated aqueous NaHCO_3 was added until gas no longer evolved. EtOAc (200 ml) and brine (200 ml) were added and the organic
- 10 phase was separated. The aqueous phase was extracted with EtOAc (150 ml, x 2). The combined organic layers were washed with brine (200 ml), dried over Na_2SO_4 , filtered and concentrated to give a crude orange oil, which was further purified on the Biotage column, eluting with hexane/EtOAc (4:1) to give **57** (4.11 g, 11.26 mmol, 68.5% yield). Electrospray MS $[\text{M}+1]^+$ 366.9.

15 Step 4:

- CH_2Cl_2 (100 ml) was added to a solution of alcohol **57** (4.11 g, 11.25 mmol) in warm DMSO (11 ml). The solution was cooled to 0 °C and P_2O_5 (6.39 g, 22.5 mmol) was then added portionwise over 30 min, followed by addition of Et_3N (6.3 ml, 45
- 20 mmol). The reaction mixture was stirred at RT for 18 h, then quenched with saturated aqueous NaHCO_3 . The aqueous layer was extracted with EtOAc (200 ml, x 3). The combined organic phases were washed with brine (200 ml), dried over Na_2SO_4 , filtered and concentrated. The crude product was triturated with hexane to give a light tan powder as desired product. The filtrate was further purified on the Biotage
- 25 column, eluting with hexane/EtOAc (4:1) to give **58** as tan solid (2.46 g, 6.74 mmol, 60.25%). Electrospray MS $[\text{M}+1]^+$ 364.1.

Step 5:

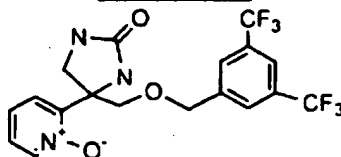
- 43 -

A suspension of ketone **58** (2.30 g, 6.30 mmol) in a mixture of (1:1) water/EtOAc (30 ml) was treated with $(\text{NH}_4)_2\text{CO}_3$ (2.12 g, 22.1 mmol) followed by KCN (0.62 g, 9.5 mmol). The solution was heated to 60 °C and stirred overnight. After cooling to RT, the reaction mixture was taken up in EtOAc (100 ml) and washed with water (100ml x 2). The separated organic layer was dried over Na_2SO_4 , filtered and concentrated *in vacuo*. The crude product was purified on a Biotage column, eluting with hexane/EtOAc (1:1) to give the desired product **59** as a white solid (2.03 g, 4.69 mmol, 74.5% yield). Electrospray MS $[\text{M}+1]^+$ 434.1.

Step 6:

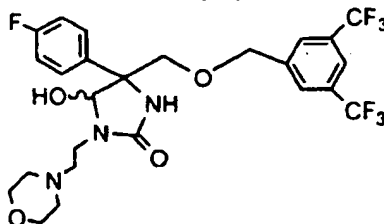
To a cooled flask containing AlCl_3 (614 mg, 4.6 mmol) at 0 °C, a 1M solution of LAH in Et_2O (3.5 ml, 3.5 mmol) was added dropwise. 10 min later, a solution of hydantoin **59** (500 mg, 1.15 mmol) in anhydrous THF (10 ml) was added slowly. After 25 min at 0 °C, the reaction was complete as determined by TLC. Water (3 ml) was added slowly to the reaction mixture, followed by 15% NaOH (3 ml) and water (9 ml). After stirring at RT for 15 min, the emulsion was filtered and rinsed with EtOAc and THF. The filtrate was washed with water (50 ml), dried over Na_2SO_4 , filtered and concentrated. The crude product was purified on the Biotage column, eluting with EtOAc/ Et_3N (9:1) to give the title compound as a solid (0.229 g, 0.546 mmol, 47.5 % yield). Electrospray MS $[\text{M}+1]^+$ 420.1.

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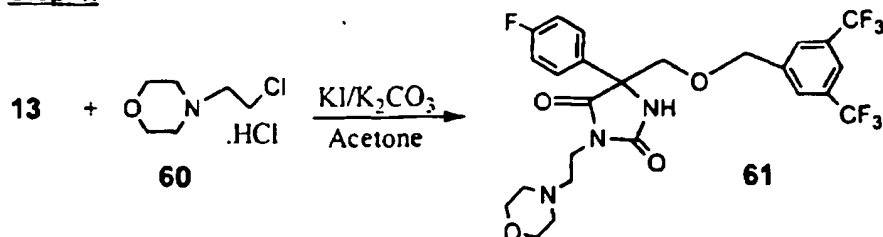
Example 24

The title compound was prepared by a similar method to that described in Example 23, Step 2, using the cyclic urea, **56**, of Example 23 in place of ether **55**. Yield was 67.1%. Electrospray MS $[\text{M}+1]^+$ 436.1.

25

Example 25

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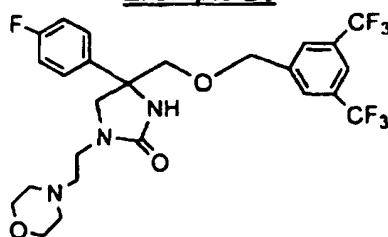
Step 1:

To the flask containing compound **13** (1.35 g, 3 mmol) in acetone (45 ml), 4-(2-chloroethyl)morpholine hydrochloride salt, **60**, (0.614 g, 3.3 mmol) was added, followed by KI (0.1 g, 0.6 mmol) and K₂CO₃ (1.24 g, 9 mmol). The reaction mixture was heated to reflux for 44 h. After cooling to RT, EtOAc (60 ml) was added and the organic mixture was washed with water (20 ml), dried over MgSO₄, filtered and concentrated to give **61** as an oil. The crude product was further purified on a silica gel column, eluting with 20% EtOAc/CH₂Cl₂ to give **61** (1.3g, 2.3 mmol, 76% of yield).

Step 2:

To a cooled flask containing AlCl₃ (0.35 g, 2.63 mmol) at 0 °C under a N₂ atmosphere, 1 M solution of LAH in Et₂O (2.0 ml, 2.0 mmol) was added dropwise. Stirred for 5 min, a solution of **61** (0.37 g, 0.66 mmol) in anhydrous THF (10 ml) was added slowly. The reaction mixture was allowed to stir from 0 °C to RT over the weekend and quenched with 20% NaOH (1 ml). The solids were filtered and rinsed well with CH₂Cl₂ after stirring for 15 min. The filtrate was concentrated and purified on flash grade silica gel, eluting with 5% CH₃OH in CH₂Cl₂ to give the title compound as a white foam (70 mg, 0.13 mmol, 20% of yield). High Resolution MS: [M+1]⁺ measured 566.1899, calculated 566.1890.

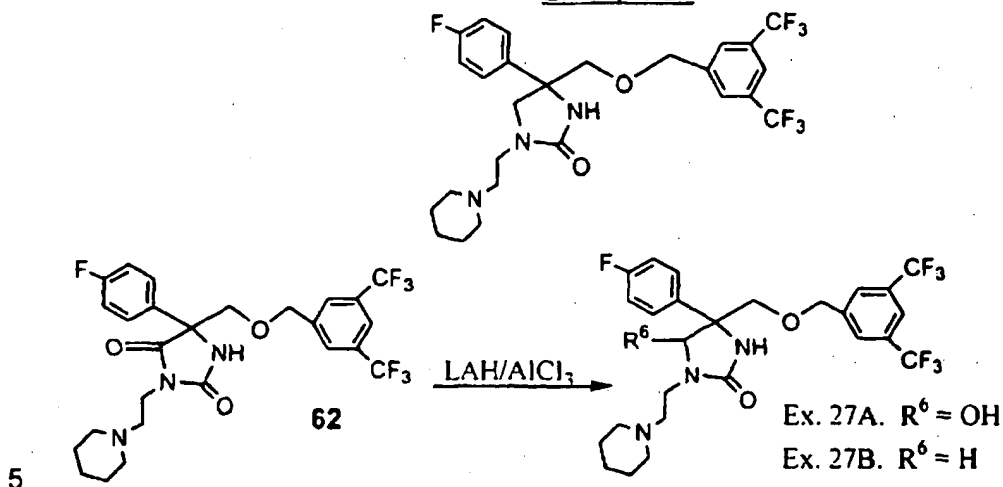
20

Example 26

To a cooled flask containing AlCl₃ (1.23 g, 9.24 mmol) at 0 °C under N₂ atmosphere, 1 M solution of LAH in Et₂O (7.0 ml, 7.0 mmol) was added dropwise. Stirred for 5 min, a solution of **61** (1.3 g, 2.3 mmol) in anhydrous THF (25 ml) was added slowly. The reaction mixture was allowed to stir at 0 °C for 15 min then heated at 70 °C for 64 h. The reaction was quenched with 20% NaOH (2 ml) and stirred at RT for 15 min. The solids were filtered and rinsed well with CH₂Cl₂. The filtrate was concentrated and purified on flash grade silica gel, eluting with 5% CH₃OH in CH₂Cl₂.

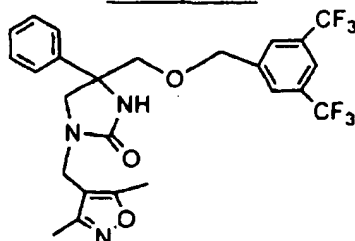
- 45 -

to give the title compound as a gummy solid (0.46g, 0.8 mmol, 36% yield). High Resolution MS: $[M+1]^+$ measured 550.1948, calculated 550.1941.

Example 27

Compound **62** was prepared in a similar method as described in Example 25 using 2-chlorethyl-1-piperidine in place of 4-(2-chloroethyl)-morpholine. A mixture of Ex. 27A and Ex. 27B was separated by flash column chromatography, eluting with 2~5% CH₃OH in CH₂Cl₂ to afford

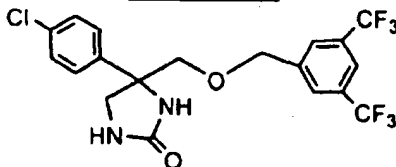
10 Ex. 27A (33% yield) (FAB MS $[M+1]^+$ 564) and Ex. 27B (43% yield) FAB MS $[M+1]^+$ 549.

Example 28

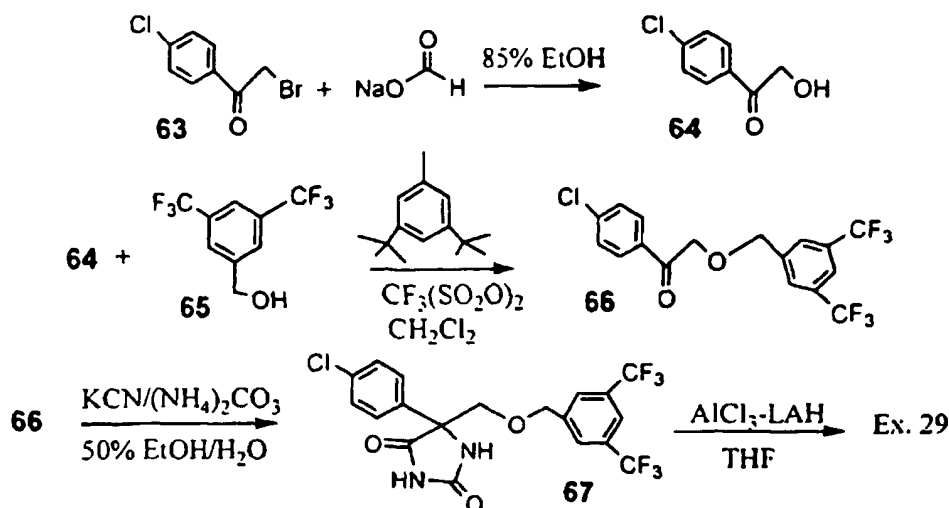
The title compound was prepared by an analogous method to that described in

15 Example 16 using the compound of Example 1 in place of the compound of Example 11 to give the title compound in a 22 % yield.

FAB MS $[M+1]^+$ 546.3.

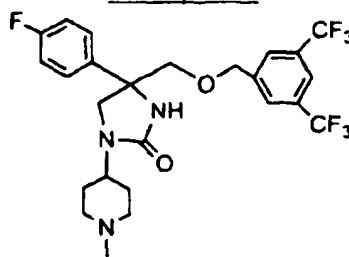
Example 29

- 46 -



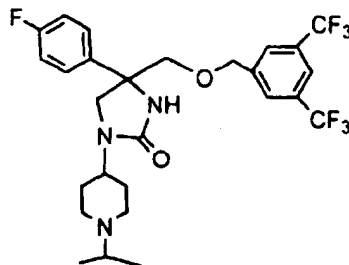
- The title compound was prepared by analogous methods as described for
- 5 Examples 1 and 2, except for the preparation of ketone intermediate **66**. Compound **66** was prepared by treatment of compound **64** (24.1 g, 98.8 mmol) and 2, 6-di-*t*-butyl-4-methyl pyridine (42 g, 206 mmol) in dry CH_2Cl_2 (400 ml) at 0 °C with slow addition of trifluoromethyl-methane sulfonic anhydride (30 g, 107 mmol) under a N_2 atmosphere. After stirring at RT for 1 h, a solution of compound **64** (14 g, 82.3 mmol)
- 10 in CH_2Cl_2 (200 ml) was added dropwise. After stirring at RT for 6 days, the solid was filtered and filtrate was washed with brine, dried (MgSO_4), filtered and concentrated to give a crude **66** as a brown oil. Compound **66** was obtained by purification through a flash grade silica gel, eluting with 8% EtOAc/hexane to give **66** as an oil (19.5 g, 50 mmol, 61 % of yield). Compound **67** was synthesized from compound **66** with a 60%
- 15 of yield and title compound was prepared from **67** by selective reduction with AlCl_3 -LAH in a 52 % yield. FAB MS $[\text{M}+1]^+$ 467.

Example 30

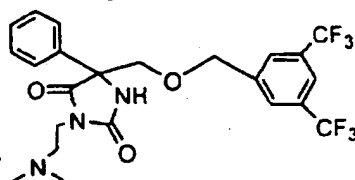


- The title compound was prepared by a method analogous to that described in
- 20 Example 7 by using compound **22** and replacing tetrahydro-4H-pyranone with 1-methyl-4-piperidone. HRMS $[\text{M}+1]^+$ calculated, 534.1992; found, 534.1987.

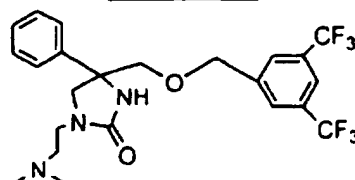
- 47 -

Example 31

- To a solution of the product of Example 10 (0.2g, 0.36 mmol) in dichloroethane (3 ml) and acetone (21mg, 0.36 mmole) at RT was added $\text{Na}(\text{OAc})_3\text{BH}$ (0.15g, 0.72 mmol). The mixture was stirred at RT for two days under a N_2 atmosphere. After completion, to the reaction was added CH_2Cl_2 (50 ml) and saturated NaHCO_3 solution, followed by routine work-up to give a crude product. Product was purified on flash grade silica gel, eluting with 3% $\text{NH}_4\text{OH}-\text{CH}_3\text{OH}$ (1:9) / 97 % CH_2Cl_2 to afford pure title compound as a solid (0.12g, 0.22 mmol) in a 62% of yield.
- 10 FAB MS $[\text{M}+1]^+$ 562.4.

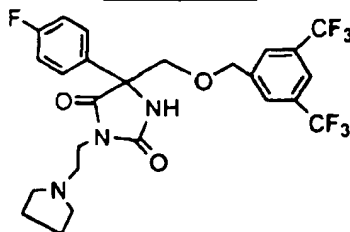
Example 32

- The title compound was prepared by a method analogous to that described in Example 25 for compound 61 in step 1, using compound 7 in place of compound 13 and N, N-dimethyaminoethyl chloride hydrochloride in place of 1-(2-chloro-ethyl)-morpholine hydrochloride. The title compound was obtained as a solid in a 96 % yield. Electrospray MS $[\text{M}+1]^+$ 504.1.
- 15

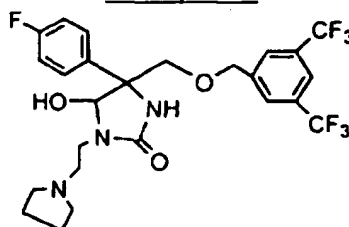
Example 33

- Reduction of the compound from Example 32 with $\text{LAH}-\text{AlCl}_3$ complex, analogous to the method described in Example 26, gave the title compound in a 32 % yield. Electrospray MS $[\text{M}+1]^+$ 490.1.
- 20

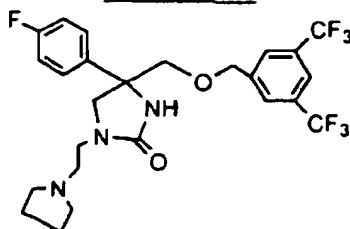
- 48 -

Example 34

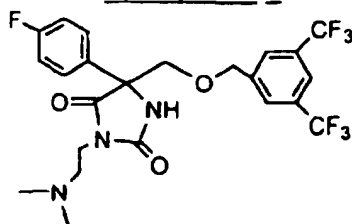
The title compound was prepared by an analogous method to that described in Example 25, Step 1, using 1-(2-chloroethyl)pyrrolidine HCl salt in place of 1-(2-chloroethyl)-morpholine hydrochloride salt. The title compound was obtained as a solid in a 77 % yield. Electrospray MS $[M+1]^+$ 548.1.

Example 35

Reduction of compound the product of Example 34 with LAH- $AlCl_3$ complex as described in Example 26, heating at 70 °C for 1 day, gave the title compound in a 36% yield. Electrospray MS $[M+1]^+$ 550.1

Example 36

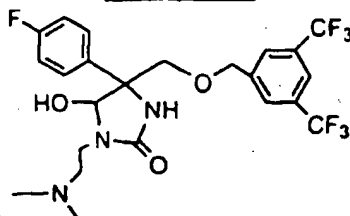
Reduction of the product of Example 34 with LAH- $AlCl_3$ complex at 70 °C for 1 day, analogous to the method as described in Example 26, gave the title compound in a 11% yield. Electrospray MS $[M+1]^+$ 534.1.

Example 37

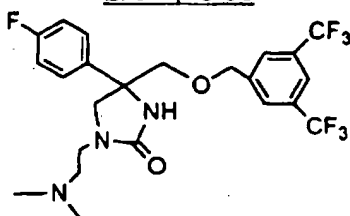
The title compound was prepared by an analogous method to that described in Example 25, Step 1, using 2-dimethylamine ethyl chloride hydrochloride salt in place

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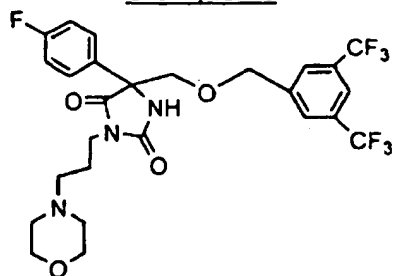
of 1-(2-chloroethyl)-morpholine hydrochloride salt. The title compound was obtained as a solid in a 83 % of yield. Electrospray MS $[M+1]^+$ 522.1.

Example 38

- 5 Reduction of the product of Example 37 with LAH- $AlCl_3$ complex at 70 °C for 1 day, analogous to the method described in Example 26, gave the title compound in a 26% yield. Electrospray MS $[M+1]^+$ 524.1

Example 39

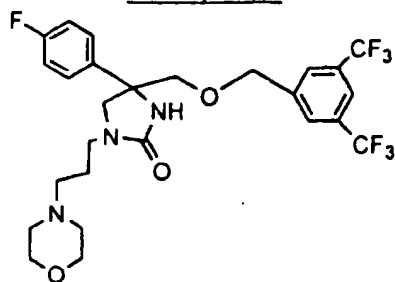
- 10 Reduction of the product of Example 38 with LAH- $AlCl_3$ complex at 70 °C for 1 day, analogous to the method described in Example 26, gave the title compound in a 15% yield. Electrospray MS $[M+1]^+$ 508.1.

Example 40

- 15 The title compound was prepared by an analogous method to that described in Example 25, Step 1, using 3-chloropropyl-morpholine in place of 1-(2-chloroethyl)-morpholine hydrochloride. The title compound was obtained as a solid in a 57 % yield. FAB MS $[M+1]^+$ 578.1.

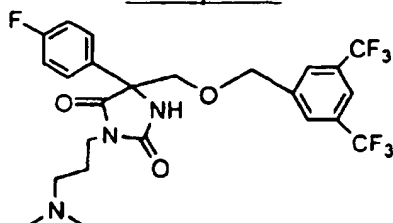
- 50 -

Example 41



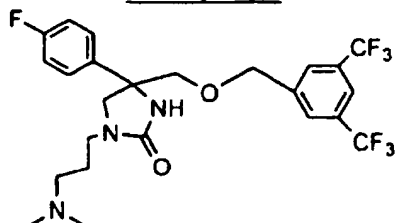
Reduction of the compound from Example 40 with LAH- AlCl_3 complex, analogous to the method as described in Example 26, gave the title compound in a 9 % yield. Electrospray MS $[\text{M}+1]^+$ 564.1.

Example 42



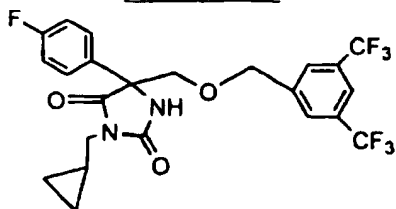
The title compound was prepared by an analogous method to that described in Example 25, Step 1, using N, N-dimethylaminopropyl chloride HCl in place of 1-(2-chloroethyl)-morpholine hydrochloride. The title compound was obtained as a solid in a 69 % yield. FAB MS $[M+1]^+$ 536.1

Example 43



Reduction of the compound from Example 42 with LAH- AlCl_3 complex,
15 analogous to the method as described in Example 26, gave the title compound in a
1.5 % yield. Electrospray MS $[\text{M}+1]^+$ 522.3.

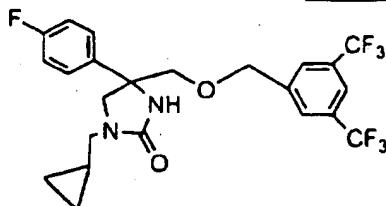
Example 44



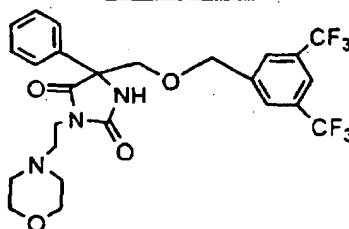
The title compound was prepared by an analogous method to that described in
20 Example 25, Step 1, using bromomethyl cyclopropane replacing 1-(2-chloroethyl)-

- 51 -

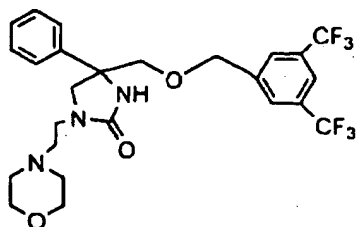
morpholine hydrochloride. The title compound was obtained as a solid in a 94 % of yield. FAB MS $[M+1]^+$ 505.1.

Example 45

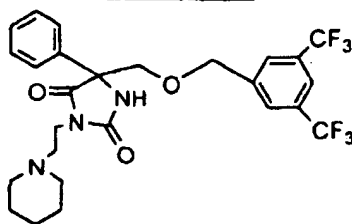
- 5 Reduction of the compound from Example 44 with LAH- $AlCl_3$ complex, analogous to the method as described in Example 26, gave the title compound in a 57 % yield. Electrospray MS $[M+1]^+$ 491.1.

Example 46

- 10 The title compound was prepared by an analogous method to that described in Example 25, Step 1, using compound 7 in place of compound 13. The title compound was obtained as a solid in a 90 % yield. FAB MS $[M+1]^+$ 546.1.

Example 47

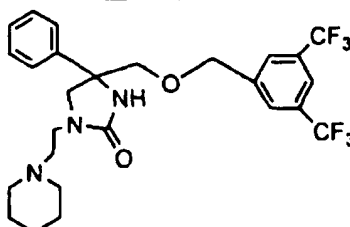
- 15 Reduction of the compound from Example 46 with LAH- $AlCl_3$ complex, analogous to the method as described in Example 26, gave the title compound in a 67 % yield. Electrospray MS $[M+1]^+$ 532.1.

Example 48

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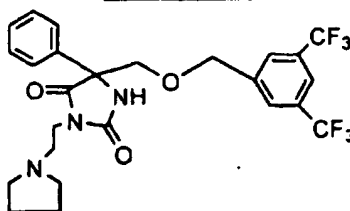
The title compound was prepared by an analogous method to that described in Example 25, Step 1, using compound **7** in place of compound **13** and 1-(2-chloroethyl)-piperidine hydrochloride in place of 1-(2-chloroethyl)-morpholine hydrochloride. The title compound was obtained as a solid in a 90 % yield. FAB MS $[M+1]^+$ 544.3.

5

Example 49

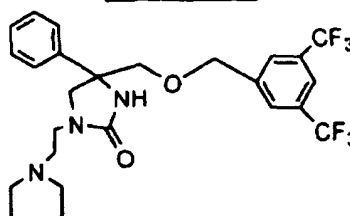
Reduction of the compound from Example 48 with LAH- $AlCl_3$ complex, analogous to the method as described for Example 26, gave the title compound in a 52 % yield. Electrospray MS $[M+1]^+$ 530.1.

10

Example 50

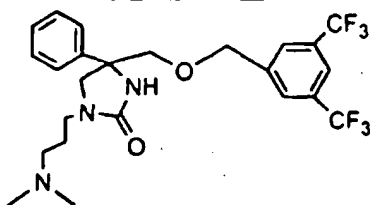
The title compound was prepared by an analogous method to that described in Example 25, Step 1, using compound **7** in place of compound **13** and 1-(2-chloroethyl)-pyrrolidine hydrochloride in place of 1-(2-chloroethyl)-morpholine hydrochloride.

15 The title compound was obtained as a solid in a 76 % yield. Electrospray MS $[M+1]^+$ 530.1.

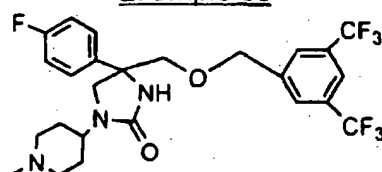
Example 51

20 Reduction of the compound from Example 50 with LAH- $AlCl_3$ complex, analogous to the method as described in Example 26, gave the title compound in a 65 % yield. Electrospray MS $[M+1]^+$ 516.1.

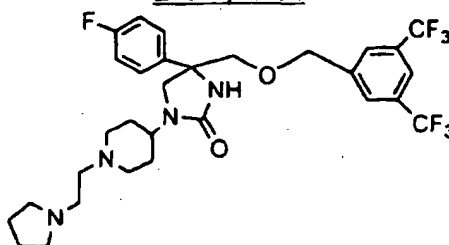
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Example 52

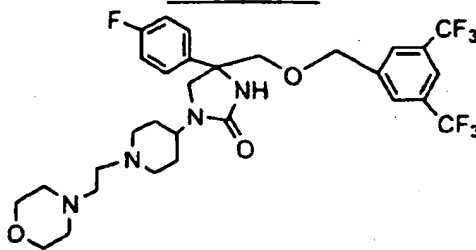
The title compound was prepared by a method analogous to Example 42, using **7** in place of **13**. Reduction of the resulting hydantoin with LAH/ AlCl_3 complex, analogous to the method described in Example 26, gave the title compound in 8 % yield. FAB MS $[\text{M}+1]$ 504.1.

Example 53

The title compound was prepared by a method analogous to that described in Example 7, using N-methylpiperidone in place of tetrahydro-4H-pyranone. The title compound was obtained as a solid in a 28 % yield. FAB MS $[\text{M}+1]$ 534.

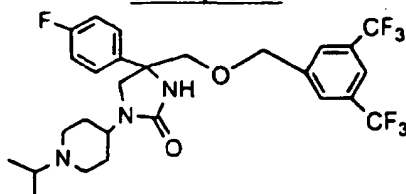
Example 54

A solution of the product of Example 10 (0.3 g, 0.54 mmol) in CH_2Cl_2 (3 ml), N, N-diisopropylethylamine (0.24 g, 1.9 mmol), and 1-(2-chloroethyl)pyrrolidine hydrochloride (92 mg, 0.54 mmol) was stirred under an atmosphere of N_2 at RT for 14 days. After work-up, a cloudy gum was obtained as a crude product which was purified by chromatography, eluting with 4.5% NH_3 - CH_3OH (1:9) / 95.5 % CH_2Cl_2 to give the title compound as an off-white solid (20 mg, 7 %). FAB MS $[\text{M}+1]$ 617.1.

Example 55

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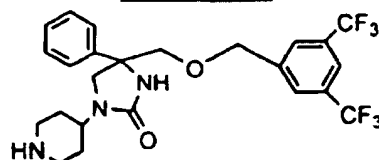
The title compound was prepared by a method analogous to Example 54, using 4-(2-chloroethyl)morpholine in place of 1-(2-chloroethyl)pyrrolidine. The title compound was obtained as a solid in a 35 % yield. FAB MS [M+1] 633.3.

Example 56

5

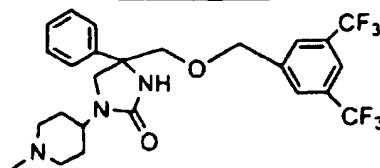
A solution of the product of Example 10 (0.2 g, 0.36 mmol) in 1, 2-dichloroethane (3 ml), acetone (21 mg, 0.36 mmol), and Na(OAc)₃BH (0.15 g, 0.72 mmol) was stirred under N₂ atmosphere at RT for 2 days. After work-up, a solid was obtained as a crude product which was then purified by chromatography, eluting with 3 % NH₃-CH₃OH (1:9) / 97 % CH₂Cl₂ to give the title compound as a white solid (120 mg, 60 %). FAB MS [M+1] 562.4.

10

Example 57

The title compound was prepared by a method analogous to Example 10, using 6 in place of 22. The title compound was obtained as a HCl salt, FAB MS [M+1] 502.1.

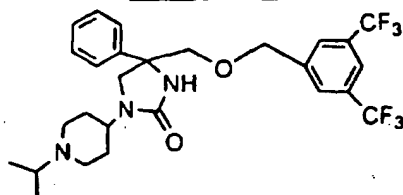
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Example 58

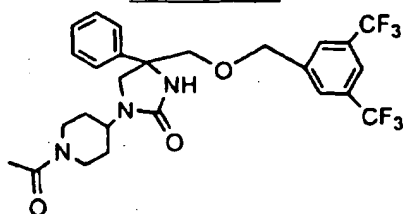
A solution of the product of Example 57 (0.3 g, 0.56 mmol) in CH₂Cl₂ (3 ml), N, N-diisopropylethylamine (0.14 g, 1.12 mmol), and CH₃I (71 mg, 0.5 mmol) was stirred under N₂ atmosphere at 0°C for 2 h. After work-up, a solid was obtained as a crude product which was then purified by chromatography, eluting with 3.5% NH₃-CH₃OH (1:9) / 96.5 % CH₂Cl₂ to give the title compound as a white solid (90 mg, 36 %). FAB MS [M+1] 516.1.

20

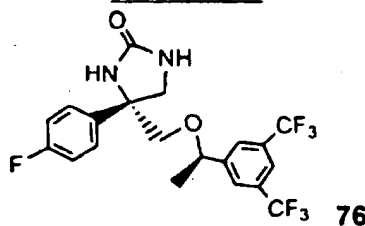
- 55 -

Example 59

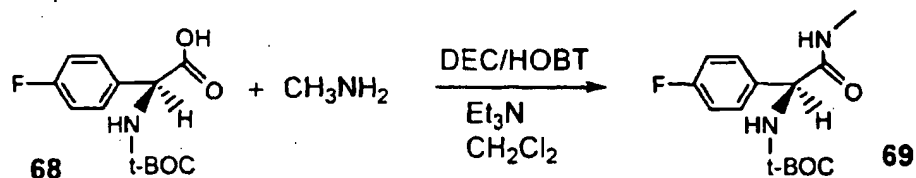
The title compound was prepared by a method analogous to Example 56, using the product of Example 57 in place of Example 10. The title compound was obtained as a solid in a 52 % yield. FAB MS [M+1] 544.2.

Example 60

The title compound was prepared by a method analogous to Example 15, using Example 57 in place of Example 14. The title compound was obtained as a solid in a 80 % yield. FAB MS [M+1] 544.1.

Example 61

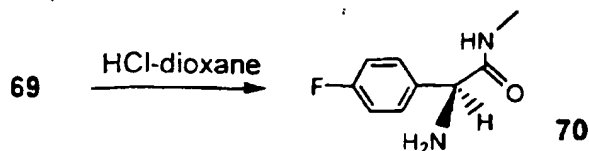
Step 1:



To a cooled solution of (S)-N-tBoc-4-fluoro-phenyl glycine in anhydrous CH_2Cl_2 at 0°C , HOBT (3.66 g, 22.4 mmol), Et_3N (3.12 ml, 22.4 mmol), DEC (4.29 g, 22.4 mmol) and 2M CH_3NH_2 in THF (10.23 ml, 20.46 mmol) were added. After stirring at 0°C for 30 min, the reaction mixture was gradually warmed up to RT under N_2 protection over night. Diluted with CH_2Cl_2 (300 ml), the reaction mixture was washed with 10% citric acid (2X100 ml), saturated NaHCO_3 -NaCl solution (2X100 ml), brine (1X100 ml). The combined organic layer was dried over MgSO_4 , filtered and concentrated *in vacuo* to give crude 69 (4.88 g).

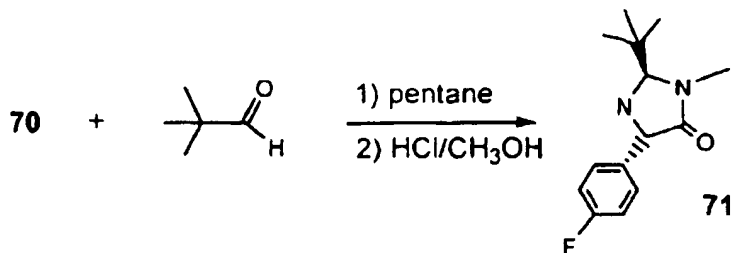
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Step 2:



- To a cooled solution of 69 (24.3 g, 86 mmol) in CH_2Cl_2 (58.8 ml) and CH_3OH (30 ml) at 0 °C was added 4M HCl-dioxane solution (200 ml, 800 mmol). The clear solution was kept at 0 °C for 2.5 hours. Solvent was evaporated and the solid material was taken up with distilled water (200 ml), neutralized with NaOH (3.4 g) in water (100 ml). Final pH was adjusted to 9.5 with K_2CO_3 . This aqueous solution was extracted with CH_2Cl_2 (4X250 ml). The combined organic layer was dried over MgSO_4 , filtered and concentrated *in vacuo* to give a white solid which further turned into a pale yellow solid 70. (14.9 g, 96%).

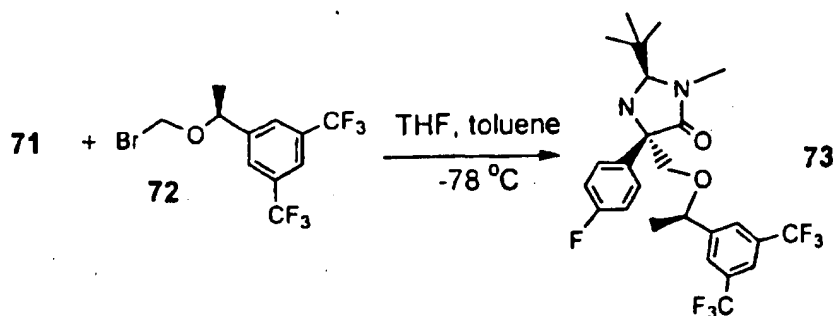
Step 3:



- A mixture of compound 70 (14.84 g, 81.5 mmol), pentane (85 ml) and pivaldehyde (11.13 ml, 101.88 mmol) was heated to 65 °C with a Dean-Stark trap for 2.5 hours under N_2 . The reaction mixture was then concentrated and the crude oil was taken up with CH_3OH (22.5 ml), cooled down to 0 °C, and the crude product was treated with saturated HCl- CH_3OH (52.5 ml) through a dropping funnel over 20 min. and stirred at ambient temperature over night. Volatile solvent was again evaporated and the residue was suspended in a mixture of brine (100 ml) and 2% K_2CO_3 solution (150 ml). This basic solution (pH=9.5) was extracted with CH_2Cl_2 (4X200 ml) and the combined organic layer was dried over MgSO_4 , filtered and concentrated *in vacuo*. The crude product (20 g) was purified on a silica gel column, eluting with 25% EtOAc in hexane (4 l), 30% EtOAc in hexane (4 l) to give compound 71 (11.67 g, 58%) as an oil.

Step 4:

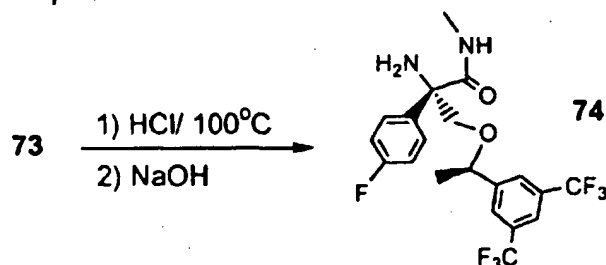
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To a cooled solution of **71** (11.5 g, 46.08 mmol) in freshly distilled degassed THF (62 ml) and anhydrous degassed toluene (192 ml) at $-78\text{ }^\circ\text{C}$, 0.5M KHMDS (97 ml, 48.5 mmol) in toluene was added slowly over 10 min through an addition funnel.

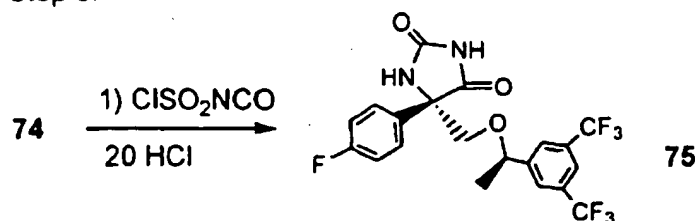
- 5 The solution was stirred at $-78\text{ }^\circ\text{C}$ for 40 min before **72** (17 g, 49 mmol) was added via a syringe. Kept stirring at this low temperature for another 2.5 hours, the reaction was quenched with saturated NH_4Cl solution (350 ml). The separated aqueous layer was extracted with CH_2Cl_2 (3X200 ml). The combined organic layer was dried over MgSO_4 , filtered and concentrated. The crude product was further purified by flash
- 10 chromatography, eluting with 15% EtOAc in hexane (6 l), 20% EtOAc in hexane (4 l). **73** was obtained as an oil (13.06 g, 56%).

Step 5:



- A mixture of **73** (12.55 g, 24.2 mmol) and 6N HCl (132 ml) was heated to
- 15 $100\text{ }^\circ\text{C}$ under N_2 for 4 h. After cooling to RT, volatile solvent was evaporated. The gummy material was redissolved in 2.5% K_2CO_3 solution (300 ml) and brine (100 ml). The aqueous solution was extracted with CH_2Cl_2 (4X250 ml). The combined organic layer was dried over MgSO_4 , filtered and concentrated *in vacuo* to give a gummy material (10.9 g) which was purified on a silica gel column, eluting with 20% EtOAc in
- 20 hexane then 1.5% $\text{NH}_3\text{-CH}_3\text{OH}$ in CH_2Cl_2 . **74** was obtained as an oil (8.45 g, 78%).

Step 6:



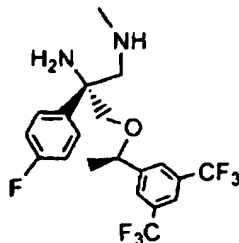
- 58 -

To a cooled solution of **74** (7.5 g, 16.6 mmol) in anhydrous CH_2Cl_2 (85 ml) at 0°C was added ClSO_2NCO (1.45 ml, 16.6 mmol) using a syringe. After stirring for 20 min, the reaction was finished and the solvent was evaporated. Taken up with 1,4-dioxane (70 ml) and 4N HCl (70 ml), the reaction mixture was then heated up to 95°C under N_2 for 4.5 hours. TLC showed completed reaction. Evaporation of volatile solvents yielded crude gummy material, which was taken up with 2.5% K_2CO_3 solution (300 ml) and CH_2Cl_2 (200 ml). The aqueous solution was extracted with CH_2Cl_2 (4X200 ml). The combined organic layer was dried over MgSO_4 , filtered and concentrated. The crude product was purified by flash chromatography, eluting with 3.5% (1:9) $\text{NH}_4\text{OH}-\text{CH}_3\text{OH}$ in CH_2Cl_2 to give **75** as a solid (6.88 g, 90%).

Step 7:

To a cooled flask containing AlCl_3 (4.3 g, 32.16 mmol) was added LAH (24.12 ml, 24.12 mmol) at 0°C under N_2 . After 15 min, a solution of **75** (3.73 g, 8.04 mmol) in anhydrous THF (126 ml) was added through a syringe. The reaction mixture was stirred at 0°C for 10 min before warming up to RT. After 4 h, the reaction was quenched with saturated Na_2SO_4 solution (10 ml) in an ice-bath. Diluted with THF (200 ml) and stirred for 1 h at RT, the reaction was dried over MgSO_4 , filtered and concentrated to give crude **76** (3.5 g), which was purified by flash chromatography eluting with 1% $\text{NH}_4\text{OH}/\text{CH}_3\text{OH}$ (1:9) in CH_2Cl_2 . mp $136-137^\circ\text{C}$, FABMS $[\text{M}+1]^+ 451.1$; $[\alpha]_D^{20} +49.4^\circ\text{C}$.

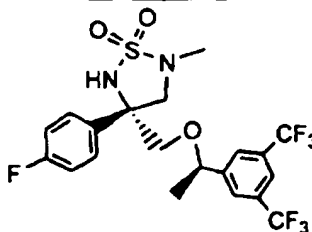
Example 62



The title compound was prepared by a method analogous to step 7 of Example 61, using compound **74** in place of compound **75**. FAB MS $[\text{M}+1]^+ 439.1$.

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Example 63

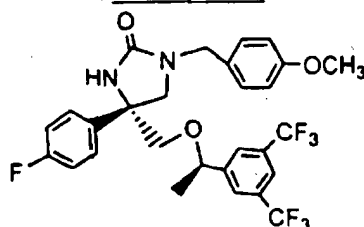


To a solution of Example **62** (180 mg, 0.41 mmol) in dioxane (4 ml), sulfamide (48.5 mg, 0.51 mmol) was added. After heating to 110°C for 5 h, the reaction was

- 59 -

concentrated. The residue was purified by flash chromatography, eluting with 4% $\text{NH}_4\text{OH}:\text{CH}_3\text{OH}(1:9)$ in CH_2Cl_2 . The title compound was collected as off-white solid (35 mg, 17%). FAB MS $[\text{M}+1]^+$ 518.1.

Example 64

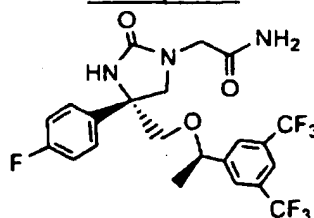


5

To a cooled solution of NaH (27 mg, 0.67 mmol) in anhydrous DMF (5 ml), was added compound 76 from Example 61 (0.25 g, 0.56 mmol) at 0 °C. After 15 min, the cold bath was removed. The reaction was warmed to RT for 45 min before p-methoxy benzyl-chloride (0.08 ml, 0.59 mmol) was added. The reaction was then stirred at RT for another 2 h. Water was used to quench the reaction and the aqueous solution was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered and concentrated. Crude material was purified through a silica gel column, eluting with 2:1 hexane to EtOAc to give the title compound as a solid (86.8 mg, 27%). FAB MS $[\text{M}+1]^+$ 571.1.

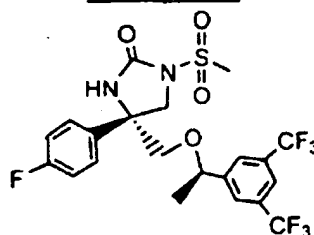
15

Example 65



The title compound was prepared by a method analogous to Example 64, using bromo acetamide in place of p-methoxy benzyl chloride. FAB MS $[\text{M}+1]^+$ 508.1.

Example 66

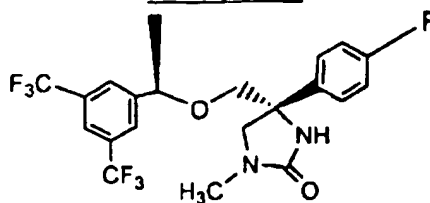


20

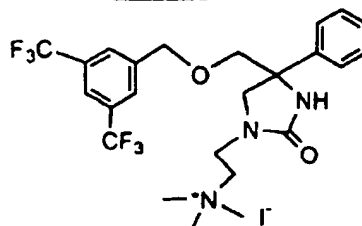
To a cooled solution of 76 (0.2 g, 0.44 mmol) in anhydrous CH_2Cl_2 (4.2 ml) at 0 °C, diethylisopropylamine (0.12 ml, 0.69 mmol) was added dropwise followed by $\text{CH}_3\text{SO}_2\text{Cl}$ (0.04 ml, 0.52 mmol). The reaction was warmed to RT and stirred overnight. The volatile solvent was evaporated and the crude product was purified on

- 60 -

a silica gel column, eluting with 4% CH₃OH (saturated with NH₃) in CH₂Cl₂. FAB MS [M+1]⁺ 529.1.

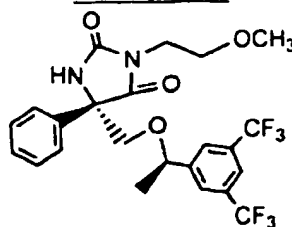
Example 67

- 5 The title compound was prepared by methods analogous to Example 1, method 1, step 5, using the compound of Example 62 in place of 6. The title compound was obtained as a solid in a 51% yield. Electrospray MS [M+1]⁺ 465.1.

Example 68

- 10 To a solution of compound of Example 67 (0.31g, 0.64 mmol) in anhydrous THF (12.8 ml) at RT, CH₃I (0.09 ml, 1.28 mmol) was added. The reaction mixture was stirred under N₂ for 5 h. Evaporation of the solvent gave a yellow foam, which was further purified by flash chromatography eluting with 7.5% CH₃OH (saturated with NH₃) in CH₂Cl₂. Electrospray MS [M+1]⁺ 504.

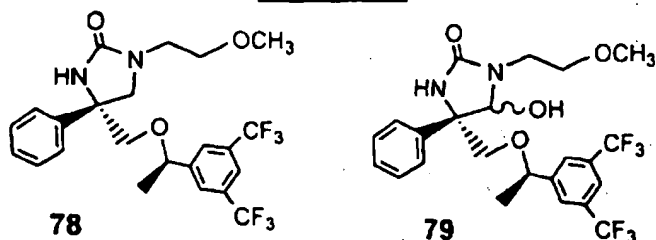
15

Example 69

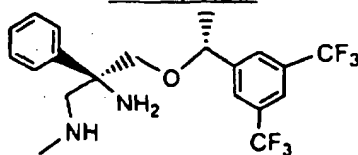
The title compound was prepared by a method analogous to Example 25, step 1, using 75 in place of 13 and 2-bromoethyl methyl ether in place of 4-(2-chloroethyl)-morpholine hydrochloride salt. The title compound was obtained in a 61% yield.

- 20 Electrospray MS [M+1]⁺ 505.1.

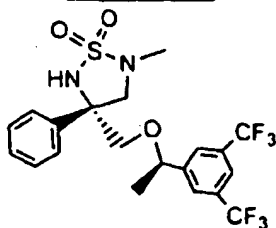
- 61 -

Example 70

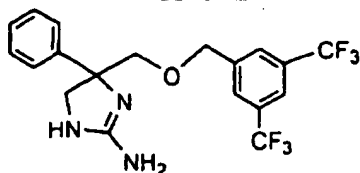
- Reduction of the product of Example 69 with LAH- AlCl_3 complex at RT overnight, analogous to the method described in Example 26, gave the title compound **78** as a clear oil and compound **79** as a crystalline solid after separation of crude product using 3% $\text{CH}_3\text{OH}(\text{NH}_3)$ in CH_2Cl_2 . Electrospray MS: compound **78**, MS $[\text{M}+1]^+$ 491.1 and compound **79**, MS $[\text{M}+1]^+$ 507.4.

Example 71

- The title compound was prepared by analogous methods to that described in Examples 61, Step 7 and 62, using the phenyl analog in place of **74** in Example 62. The title compound was obtained as an oil. MS $[\text{M}+1]^+$ 407.1.

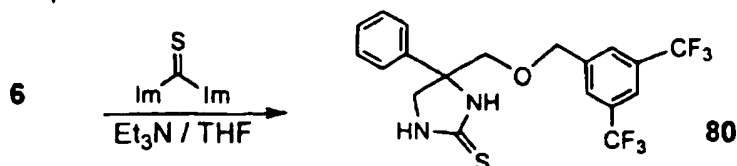
Example 72

- The title compound was prepared in a method analogous to Example 63, using Example 71 in place of the compound of Example 62. The title compound was obtained as an oil in 44% yield. NMR (CDCl_3) 1.45 (d, 3H), 2.38 (s, 3H), 3.87 (d, 1H), 3.27 (d, 1H), 3.45 (d, 1H), 3.55 (d, 1H), 4.55 (q, 3H), and aromatic protons (7.3-7.4, 7.58, 7.78).

Example 73**82**

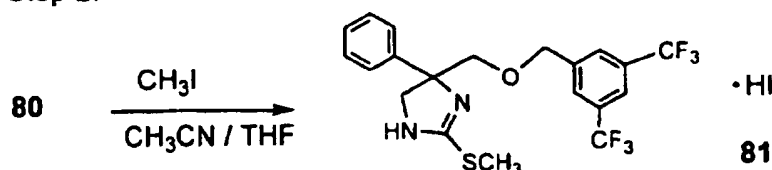
- 62 -

Step 1.



To a solution of compound 6 (1.85 g, 3.97 mmol) in dry THF (30 ml) was added Et₃N (0.6 ml, 4.3 mmol), 4 Å molecular sieve (3 g) and thiocarbonyl diimidazole (1.179 g, 5.95 mmol). The mixture was stirred at 0 °C under N₂ for 20 h. After completion, molecular sieve was filtered off and THF evaporated. The residue was redissolved in EtOAc (300 ml), washed with 0.04 N HCl (50 ml, 2X), dried (MgSO₄), filtered and evaporated. Pure 80 was obtained as a solid (1.41 g, 3.25 mmol) in 82% yield by flash silica chromatography (50 g), eluting with 5 % (1:9) [NH₄OH/CH₃OH] / 95 % CH₂Cl₂.

Step 2.

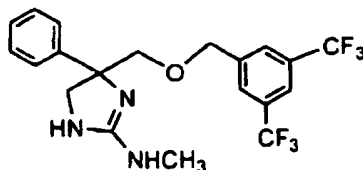


To a solution of 80 (0.75 g, 1.73 mmol) in CH₃CN and THF (6 ml) was added CH₃I (0.13 ml, 2.1 mmol). The solution was stirred at RT overnight under an atmosphere of N₂. After completion, solvent was evaporated and the product used in the next step without further purification.

Step 3

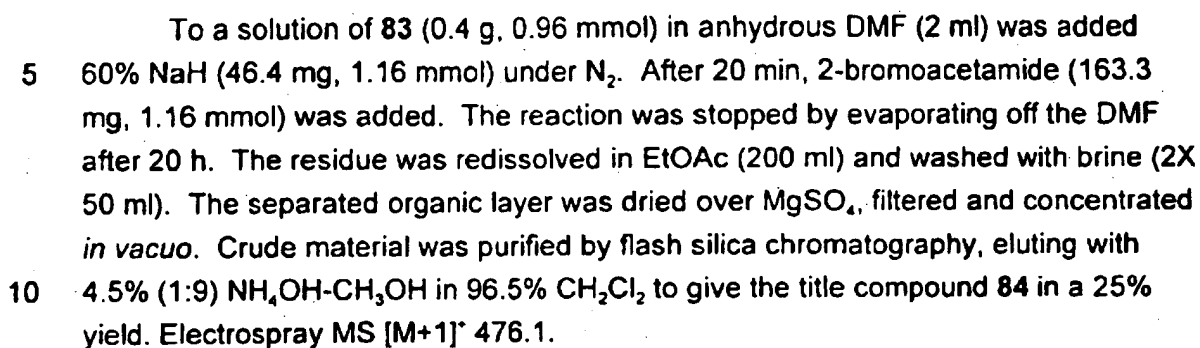
81 (150 mg, 0.334 mmol) was treated with 2 M NH₃-CH₃OH solution (1 ml). The solution was stirred in a vial with a sealed cap for 5 days. After completion, solvent was evaporated and product was purified by flash silica gel (50 g), eluting with 5 % (1:9) NH₄OH- CH₃OH/ 95 % CH₂Cl₂. LCMS [M+1]⁺ 418.1.

Example 74



The title compound was prepared in a method analogous to Example 73, step 3, using 2M NH₂CH₃ in place of 2M NH₃. Product was purified on flash silica gel (50 g), eluting with 5 % (1:9) NH₄OH- CH₃OH/ 95 % CH₂Cl₂. LCMS [M+1]⁺ 432.1.

Example 75

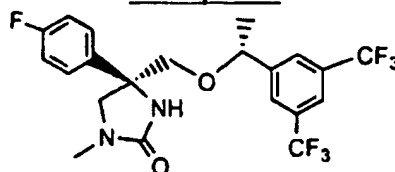


85

To a flask containing Example 71 (0.2 g, 0.48 mmol) and molecular sieves (0.18 g) in anhydrous toluene (4.5 ml), glacial acetic acid (0.3 ml, 5.24 mmol) was added. The reaction mixture was heated to 100 °C for 48 h. Cooled down to RT, the reaction was worked up by pouring into a separatory funnel containing EtOAc and saturated NaHCO₃. The aqueous layer was extracted with EtOAc (2X 50 ml). The combined organic layer was washed with brine (50 ml), dried over Na₂SO₄, filtered and concentrated. The title compound **85** was obtained by purifying through flash chromatography, eluting with 20% EtOAc in hexane. Yield: 43%. Electrospray MS [M+1]⁺ 445.1.

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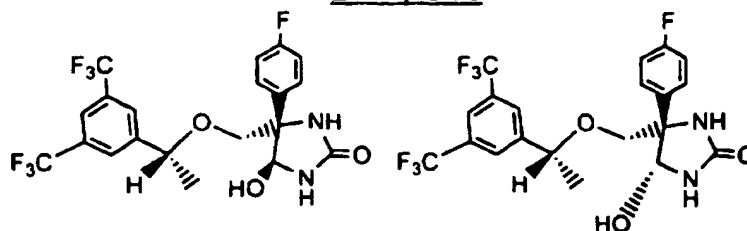
The title compound was prepared in a method analogous to Example 76, using propanoic acid in place of glacial acetic acid. The title compound was obtained in a 25% yield. Electrospray MS $[M+1]^+$ 459.1.

Example 78

5

The title compound was prepared by a method analogous to Example 1, Method 1, Step 5, using Example 62 in place of 6. Purification by flash silica chromatography, using 5% (1:9)[NH₄OH/ CH₃OH]/95 % CH₂Cl₂ gave the title compound as a solid with a yield of 51%. LCMS $[M+1]^+$ 465.1.

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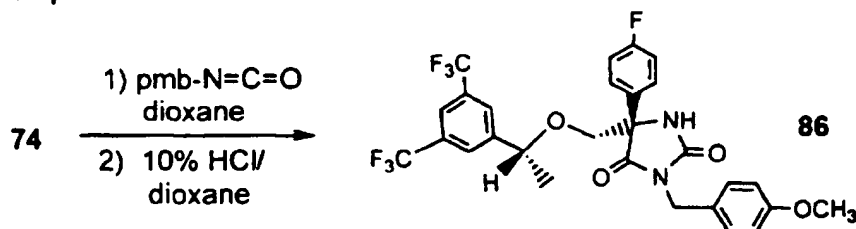
Example 79

Ex. 79a

Ex. 79b

88

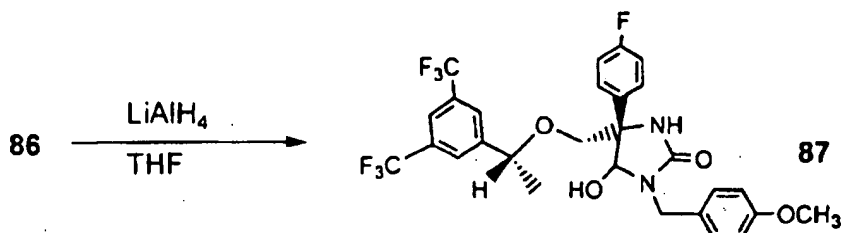
Step 1:



A solution of **74** (1.8 g, 4 mmol) in 1,4-dioxane (17 ml) and 4-methoxybenzylisocyanate (98 mg, 6 mmol) was stirred under N₂ at RT for 4 h. The mixture was mixed with 10% aqueous HCl in dioxane (17 ml), then heated to 90 °C for 5.5 h. After work-up, a solid was obtained as a crude product which was then purified by chromatography, eluting with 15% EtOAc/85% hexane to give **86** as a white solid (1.8g, 82 %). FAB MS $[M+1]^+$ 584.

20 Step 2:

- 65 -

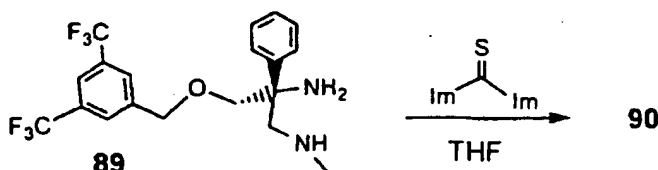
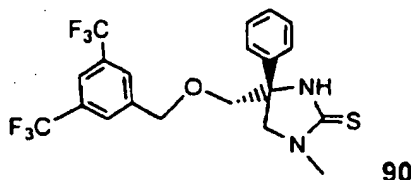


A mixture of **86** (1.7 g, 2.91 mmol) in THF (10 ml) and LiAlH_4 (1M in ether, 3.3 ml, 3.2 mmol) was stirred under N_2 at 0°C for 30 min., then RT for 3 h. After work-up, a solid was obtained as a crude product which was then purified by chromatography, eluting with 2 % $\text{NH}_3\text{-CH}_3\text{OH}$ (1:9) / 98 % CH_2Cl_2 to give **87** as a white solid (1.54g, 91 %). FAB MS $[\text{M}+1]$ 587.4.

Step 3:

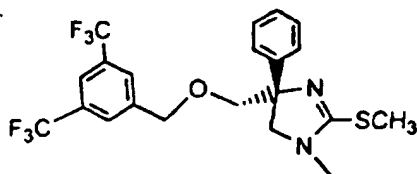
A mixture of **87** (1.48 g, 2.53 mmol) in CH_3CN (33 ml)/ H_2O (10 ml) and $\text{Ce}(\text{NH}_4)_2(\text{NO}_3)_6$ (5.54 g, 10.1 mmol) was stirred under N_2 at RT for 50 min. After work-up, an oil was obtained as a crude product which was then purified by chiral HPLC (Chiralpak[®] AD), to give the title compounds, examples 79a and 79b, as a white solid (0.19g, 16 %). FAB MS $[\text{M}+1]$ 467.1.

Example 80

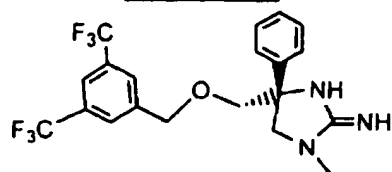


89 was prepared by a method analogous to Examples 61 and 62, using s-(+)-2-phenylglycine methyl ester hydrochloride in place of (s)-N-BOC-4-fluorophenylglycine in Example 61. A mixture of **89** (2.03 g, 5 mmol) in THF (6 ml) and 1,1'-thiocarbonyldiimidazole (1.34 g, 7.5 mmol) was stirred under N_2 at RT for 18 h. After work-up, a yellow paste was obtained as a crude product which was then purified by chromatography, eluting with 25% EtOAc/75% hexanes, to give the title compound **90** as a solid (1.35 g, 62 %). FAB MS $[\text{M}+1]$ 449.1.

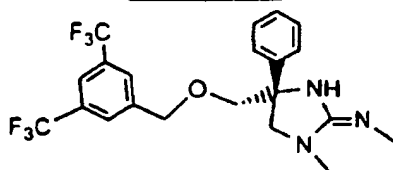
- 66 -

Example 81

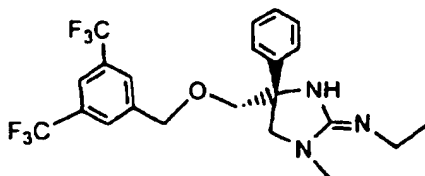
A solution of the product of Example 80 (1.2 g, 2.68 mmol) in THF (10 ml) and CH_3I (0.45 g, 3.2 mmol) was stirred under N_2 at RT for 5 h. After work-up, a solid was obtained as a crude product which was then purified by trituration with ether to give the title compound as a pale yellow salt (1.5 g, 94 %). FAB MS $[\text{M}+1]$ 463.1.

Example 82

A solution of the product of Example 81 (0.3 g, 0.51 mmol) and 7 N NH_3 (2 ml, 14 mmol) in CH_3OH was stirred under N_2 at RT for 4 days. After work-up, a solid was obtained as a crude product which was then purified by chromatography, eluting with 9 % NH_3 - CH_3OH (1:9) / 91 % CH_2Cl_2 to give the title compound as a solid (0.19 g, 86 %). FAB MS $[\text{M}+1]$ 432.1.

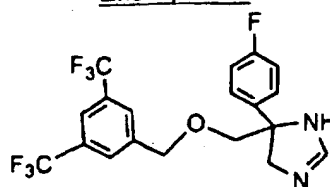
Example 83

The title compound was prepared by a method analogous to Example 82, using 2 M CH_3NH_2 in CH_3OH in place of 7 M NH_3 in CH_3OH . The title compound was obtained as a solid in a 95 % yield. FAB MS $[\text{M}+1]$ 446.1.

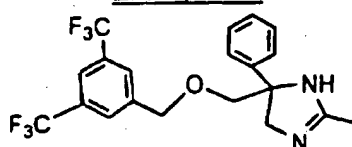
Example 84

The title compound was prepared by a method analogous to Example 82, using 2 M ethylamine in CH_3OH in place of 7 M NH_3 in CH_3OH . The title compound was obtained as a solid in a 17 % yield. FAB MS $[\text{M}+1]$ 460.1.

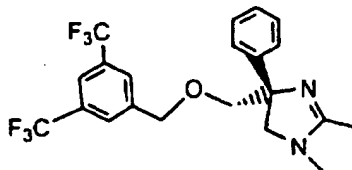
- 67 -

Example 85

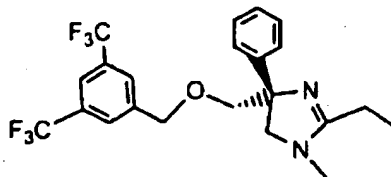
A mixture of 22 (0.3 g, 0.73 mmol) and N, N-dimethylformamide dimethyl acetal (87 mg, 0.73 mmol) was heated to 60 °C for 18 h. The reaction mixture was purified by chromatography, eluting with 3.5 % NH₃-CH₃OH (1:9)/96.5 % CH₂Cl₂ to give the title compound as an off-white solid (180 mg, 60 %). FAB MS [M+1] 421.1.

Example 86

A mixture of 6 (0.4 g, 1 mmol) in toluene (3 ml) and acetic acid (0.24 g, 4 mmol) was heated to reflux for 6 days. After work-up, an oil was obtained as a crude product which was purified by chromatography, eluting with 5 % NH₃-CH₃OH (1:9) / 95 % CH₂Cl₂ to give the title compound as a HCl salt (75 mg, 18 %) after treating the pure compound with 1 eq. of 2 N HCl-Et₂O solution. FAB MS [M+1] 417.1.

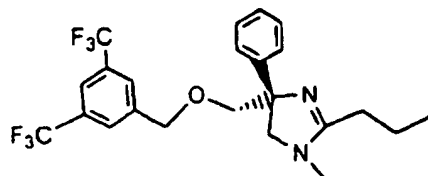
Example 87

A mixture of 89 (0.4 g, 0.985 mmol) in toluene (3 ml) and acetic acid (0.59 g, 9.85 mmol) was heated to 100 °C for 40 h. After work-up, an oil was obtained as a crude product which was purified by chromatography, eluting with 3.5 % NH₃-CH₃OH (1:9) / 96.5 % CH₂Cl₂ to give the title compound as a HCl salt (0.32 g, 76 %) after treating the pure compound with 1 eq. of 2 N HCl-Et₂O solution. FAB MS [M+1] 431.1.

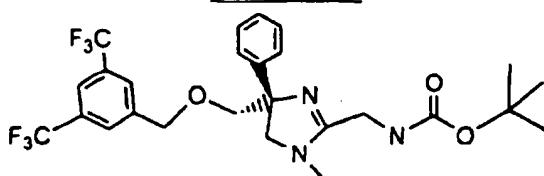
Example 88

The title compound was prepared by a method analogous to Example 87, using propionic acid in place of acetic acid. The title compound was obtained as a HCl salt in a 59 % yield. FAB MS [M+1] 445.1.

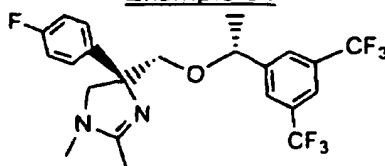
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Example 89

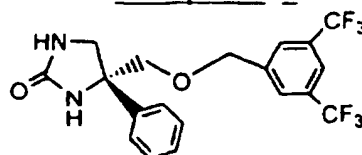
The title compound was prepared by a method analogous to Example 87, using n-butyric acid in place of acetic acid. The title compound was obtained as a
 5 HCl salt in a 88 % yield. FAB MS [M+1] 459.1.

Example 90

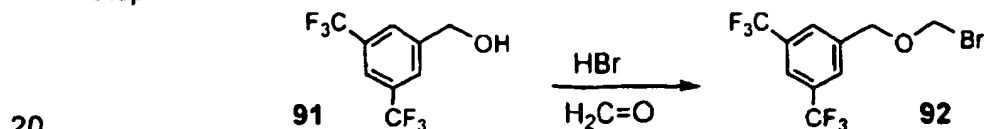
The title compound was prepared by a method analogous to Example 87, using N-t-BOC glycine in place of acetic acid. The title compound was obtained as a
 10 HCl salt in a 63 % yield. FAB MS [M+1] 546.1.

Example 91

The title compound was prepared by a method analogous to Example 76 using Example 62 in place of Example 71. Purification the crude material by flash silica
 15 chromatography, using 5% (1:9)[NH₄OH/CH₃OH]/95 % CH₂Cl₂ gave the title compound as a solid with a yield of 75%. LCMS [M+1]⁺ 463 1.

Example 92

Step 1:

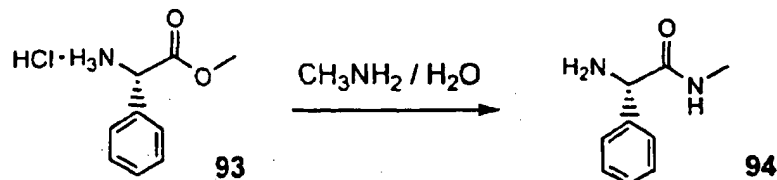


A flame-dried flask was charged with compound **91** (50 g, 204.8 mmol, 1 equiv.) and paraformaldehyde (6.76 g, 225.3 mmol, 1.1 equiv.). The solid mixture was dissolved under N₂ with a heat gun until the solution was homogeneous, approximately 20 min. The solution was allowed to cool to RT. HBr gas was bubbled

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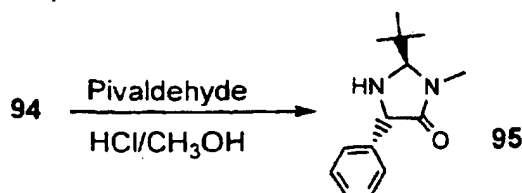
- into the solution at a fast rate at RT. The reaction was continued for 2.5 h. The layers were separated, the upper layer was diluted with hexane (150 ml) and any aqueous remnants were removed. The hexane layer was dried over MgSO_4 overnight, then filtered and concentrated. Short path distillation (high vac, 75°C) gave pure **92** (67.86 g, 98%) as a clear oil. ^1H NMR (CDCl_3) δ 4.85 (s, 2H), 5.79 (s, 2H), 7.84 (s, 2H), 7.87 (s, 1H).

Step 2:



- Compound **93** (100g, 496 mmol, 1 equiv.) was added to CH_3NH_2 (160 ml, 40% in H_2O , 4 equiv.) in a cool water bath at $10\text{--}16^\circ\text{C}$ over a period of 15 min. After complete addition, the solution was warmed to RT and stirred for 1 h. The reaction was monitored by TLC in 98:2 EtOAc/ CH_3OH . Upon completion, the reaction was quenched with brine (25% in H_2O , 500 ml). The solution was extracted with 1:1 THF/EtOAc (4 x 400 ml) and the combined organic phases were dried (Na_2SO_4) and concentrated. The residue was dried on high vacuum to give **94** (79.47 g, 98%). Electrospray MS $[\text{M}+1]^+$ 165.0.

Step 3:



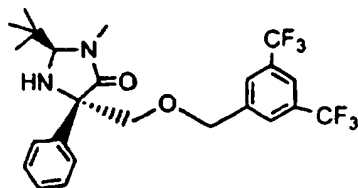
- A mixture of **94** (79.44 g, 483.8 mmol, 1 equiv.), pentane (550 ml), and trimethylacetaldehyde (65.7 ml, 604.8 mmol, 1.25 equiv.) was heated to 65°C in a system equipped with a condenser, Dean-Stark trap and N_2 inlet. The mixture was heated for 3 h and the suspension dissolved. The solution was cooled to RT and stirred overnight (16 h). The solution was concentrated, redissolved in CH_3OH (140-150 ml), cooled in an ice bath for 30 min, and then saturated $\text{HCl}-\text{CH}_3\text{OH}$ (300 ml) was slowly added via a dropping funnel over 30 min. The solution was stirred at 0°C , then warmed to RT under N_2 overnight and concentrated on high vac to yield a crude, yellow oil (109.1 g). The oil was redissolved in CH_2Cl_2 (800 ml) and washed with 25% K_2CO_3 (w/w, 400 ml). The aqueous portion was washed again with CH_2Cl_2 (2 x 400 ml). The combined organics were dried (Na_2SO_4) and concentrated to give a yellow solid (98.7 g). The crude solid was recrystallized out of hot MTBE (300 ml), then cooled to 0°C to give **95** as a white solid (54.88 g, 49%). Mother liquor still contained

- 70 -

product which could be isolated by another recrystallization or by chromatography.

Electrospray MS $[M+1]^+$ 451.1

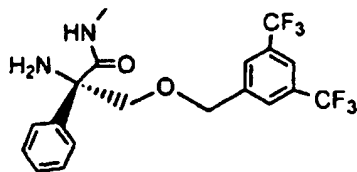
Step 4:



96

- 5 All reagents were deoxygenated under Ar prior to use. **95** (42.05 g, 181 mmol, 1 equiv.) was dissolved in dry THF from a still (550 ml) while agitating with a mechanical stirrer. The solution was cooled to approximately -70°C in dry ice/acetone and a solution of 1.5 M LDA•THF in cyclohexane (124.3 ml, 186.5 mmol, 1.03 equiv.) was added over 20 min. The resultant dark orange/brown solution was
- 10 allowed to stir at -78°C for 30 min. Bromide **92** (64.0 g, 190 mmol, 1.05 equiv.) was slowly added via syringe over 20 min. The solution was stirred at -78°C and monitored by TLC in 4:1 Hex/EtOAc. The reaction was complete after 2 h. The reaction was quenched with sat. aq. NH_4Cl (300 ml) at -78°C and then warmed to RT while stirring vigorously. Phases were separated and the organic phase washed with
- 15 H_2O (2 x 150 ml). The aqueous layer was washed with EtOAc (300 ml). The combined organics were dried (Na_2SO_4) and concentrated to give a crude, light yellow solid (32.63 g), which was recrystallized out of hot pentane to give **96** (33.66 g). The mother liquor was again concentrated to an orange solid and recrystallized out of pentane (150 ml) to give an additional 10.50 g of product (Overall yield = 44.16
- 20 g, 50%). The mother liquor (46.7 g, approximately 50% pure by NMR) still contained starting product which could be isolated. Electrospray MS $[M+1]^+$ 489.1

Step 5:



97

- To a solution of **96** (33.58 g, 68.76 mmol, 1 equiv.) in CH_3OH (300 ml),
- 25 concentrated HCl (300 ml) was added dropwise with stirring. The solution was heated to 95°C and stirred vigorously, and was stirred and heated overnight. The reaction was monitored by TLC. Upon completion, solvent was evaporated and the residue was taken up in CH_2Cl_2 (750 ml). The solution was treated with 25% aq. K_2CO_3 (approximately 350 ml) until pH 12. The mixture was filtered and the layers
- 30 separated. The organic layer was washed with 25% aq. K_2CO_3 (500 ml). Combined aqueous layers were extracted with CH_2Cl_2 (2 x 750 ml). Combined organic layers

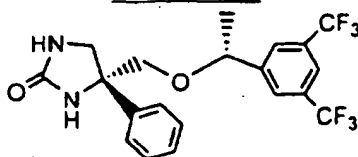
- 71 -

were dried (Na_2SO_4), and concentrated to a yellow oil. ^1H NMR shows pure **97** (29.20 g, 100%). ^1H NMR (CDCl_3) δ 2.04 (s, 2H), 2.80 (d, $J = 4.8$ Hz, 3H), 3.67 (d, $J = 8.8$ Hz, 1H), 4.53 (d, $J = 8.8$ Hz, 1H), 4.67 (d, $J = 12.5$ Hz, 1H), 4.77 (d, $J = 12.5$ Hz, 1H), 7.26-7.36 (m, 3H), 7.51 (d, $J = 8.1$ Hz, 2H), 7.65 (d, $J = 4.4$ Hz, 1H), 7.75 (s, 2H), 7.79 (s, 1H).

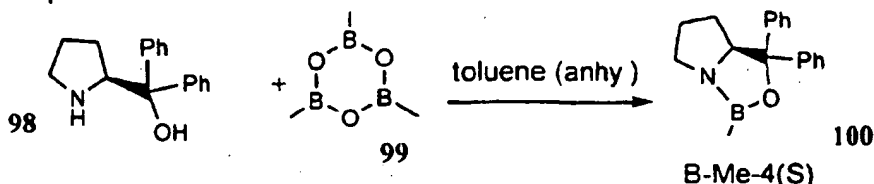
Step 6:

The title compound was prepared from **97** using procedures similar to Example 61, steps 6 and 7.

Example 93

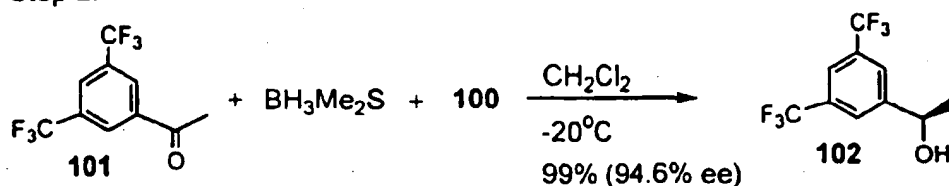


Step 1:



A flame-dried flask was charged with **98** (5.10 g, 20.1 mmol, 1.0 eq) and anhydrous toluene (56 ml). This cloudy solution was heated up to 140~150 °C. 36 ml of dry toluene was azeotropically distilled through a Dean-Stark trap with an air condenser. Another 36 ml of toluene was added. This azeotropical distillation was repeated three times to ensure **98** was totally dry. After the third azotropical distillation was done, another 36 ml of anhydrous toluene was added. The solution was allowed to cool down to RT. **99** (1.90 ml, 13.5 mmol, 0.67 eq) was syringed in within 5 min. White solid was formed at about 6 min after completion of the addition. The reaction mixture was stirred at RT for 30 min, then 36 ml of toluene was distilled off. Another 36 ml of dry toluene was added and distilled off again. The distillation was repeated one more time, and 20 ml of 1.0 M of **100** (CBS catalyst B-Me-4(S) solution in toluene) was prepared. The almost colorless solution can be used in CBS reduction directly.

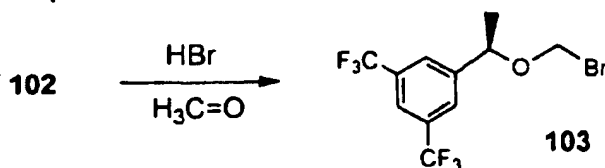
Step 2:



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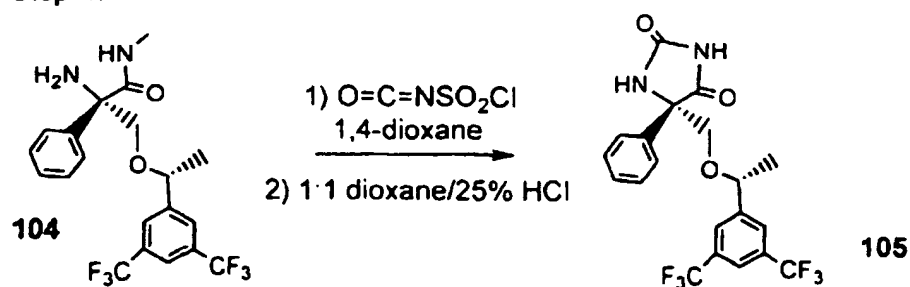
101 (102.14 g, 0.4 mol, 1.0 eq) was dissolved in anhydrous CH_2Cl_2 (780 ml) and the resulting solution was transferred into a dry dropping funnel. An oven-dried 3L flask was cooled to -20°C , and 20 ml of **102** toluene solution was syringed in, followed by 40 ml of 10.0~10.3 M borane-methylsulfide complex. Then the **101** solution was added dropwise through the dropping funnel over 2 days. During the addition, the temperature was maintained at -20°C . Once the addition was finished, the reaction was monitored by TLC in 4:1 hexane/EtOAc. When **101** was completely consumed, 250 ml of CH_3OH was added slowly, and hydrogen gas was emitted. The reaction solution was then concentrated to give white solid. The solid was dissolved in Et_2O (500 ml), then 45 ml of 2.0 M of HCl in Et_2O was added slowly at -20°C . White precipitate was formed. The reaction mixture was warmed to RT and stirred for 30~40 min. The mixture was filtered and the filtrate concentrated to give 101.5 g of white solid **102** (yield 98.7%). Chiral HPLC Chiral OD(Chiralcel) column (Hexane/IPA = 98/2) showed 94.6%ee.

Step 3:



103 was prepared in a method analogous to Example 92, Step 1 using **102** in place of **91**.

Step 4:



Amino amide **104** (14.14 g, 32.55 mmol, 1 equiv.) (prepared in a similar procedure to that described in Example 92, substituting **103** in place of **92** in step 4) was taken up in dry CH_2Cl_2 (120 ml). The solution was cooled to -78°C and chlorosulfonyl isocyanate (2.84 ml, 32.55 mmol, 1 equiv.) was added. The reaction was stirred at 0°C for 3 h and then concentrated to a white solid. The solid was dissolved in 1,4-dioxane (120 ml) and 3 N aqueous HCl (120 ml), then stirred at 90°C for 5 h, then at RT overnight. The solution was diluted with H_2O (250 ml) and extracted with EtOAc (3 x 400 ml). The combined organics were dried (Na_2SO_4), and concentrated to a crude, white foam (15.80 g). the foam was purified by plug

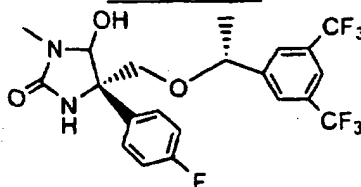
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chromatography on a 600 ml fritted funnel, eluting with 2:1 Hex/EtOAc. The fractions 2-8 were collected and concentrated to give pure **105** (10.26 g, 71%). FAB MS $[M+1]^+$ 447.1

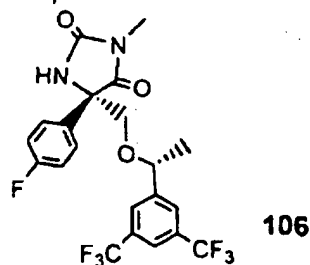
Step 5:

- 5 $AlCl_3$ (12.2 g, 91.4 mmol, 4 equiv.) was added to a flask equipped with a N_2 inlet. The flask was cooled to $0^\circ C$ and a solution of 1 M LAH in ether (68.6 ml, 68.6 mmol, 3 equiv.) was slowly added. The resultant white slurry was stirred at $0^\circ C$ for 15 min. Next, a solution of hydantoin **105** (10.2 g, 22.85 mmol, 1 equiv.) in 150 ml dry THF was added via canula. The solution was warmed to RT and stirred for 41 h. The solution was again cooled to $0^\circ C$ and H_2O (20 ml) was added, then 15% aq. NaOH (w/w, 20 ml) followed by H_2O (60 ml). The biphasic solution was stirred for 30 min. All emulsion formed was dissolved with 1 N HCl (approximately 300-400 ml) and the layers were separated. The aqueous layer was extracted with EtOAc (2 x 500 ml). The combined organics were washed with H_2O (200 ml), dried (Na_2SO_4), and
- 15 concentrated to give the crude (9.86 g). The crude material was initially purified by plug chromatography on a 2L fritted funnel, eluting with 1:1 Hex/EtOAc, followed by 98:2 EtOAc to give 8.0 g of material, which still contained 3% of a less polar impurity. The solid was recrystallized from hot MTBE (30 ml) to provide pure Example 93 (6.5 g, 79%). FAB MS $[M+1]^+$ 433.1

20

Example 94

Step 1:



- 25 **104** (3.0 g, 6.63 mmol, 1 equiv.) (prepared analogously to the procedure described in a Example 92) was dissolved in CH_2Cl_2 (100 ml) and cooled to $0^\circ C$. Et_3N (2.78 ml, 19.89 mmol, 3 equiv.) followed by triphosgene (787 mg, 2.65 mmol, 0.4 equiv.) was added and the solution was allowed to stir and warm to RT. Reaction was complete after 2 h as determined by TLC. The reaction was quenched with sat. aq. $NaHCO_3$ (100 ml) and the layers were separated. The organic layer was washed

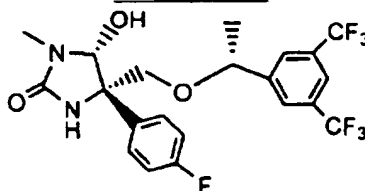
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with brine (100 ml), dried (Na_2SO_4), and concentrated. The crude product was purified by Biotage chromatography, eluting with CH_2Cl_2 to give 2.024 g (64%) of **106**. FAB MS $[\text{M}+1]^+$ 479.1.

Step 2:

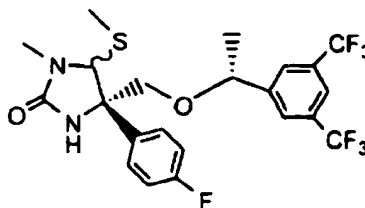
- 5 **106** (919 mg, 1.92 mmol, 1 equiv.) was dissolved in dry THF (10 ml) and cooled to 0°C . A 1 M solution of LAH in Et_2O (1.92 ml, 1.92 mmol, 1 equiv.) was slowly added and the solution was stirred at 0°C for 10 min. The reaction was warmed to RT and completed after 2 h. The solution was cooled to 0°C and water (3 ml) was slowly added, followed by 15% aq. NaOH (3 ml), and then more water (10 ml). The emulsion was dissolved with 1 M HCl and the layers were separated. The aqueous layer was extracted with EtOAc (2 x 100 ml) and the organic layers were combined, washed with water (50 ml), dried (Na_2SO_4), and concentrated to give crude product (914 mg). Biotage chromatography, eluting with 98:2 $\text{CH}_2\text{Cl}_2/\text{CH}_3\text{OH}(\text{NH}_3)$ gave a 3:2 mixture of diastereomers (798 mg) (87%). Prep HPLC on 50 mg of product using a chiralcel OD column, eluting with 85:15 Hex/IPA gave the title compound ($t_R = 6.4$ min), a white powder (12.7 mg, 51%, 98% de). HRMS $[\text{M}+1]^+$ 481.1362.

Example 95



- 20 The title compound was prepared from **106** using the procedure of Example 94. Prep HPLC on 50 mg of material on chiralcel OD column, eluting with 85:15 Hex/IPA gave the title compound ($t_R = 8.2$ min) as a clear oil (17.5 mg, 70%, 78% de). HRMS $[\text{M}+1]^+$ 481.1362

Example 96

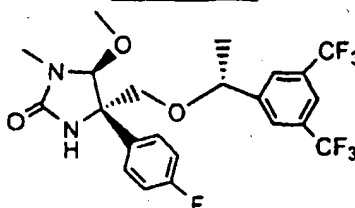


- 25 A 2:3 diastereomeric mixture of Examples 94 and 95 (100 mg, 0.208 mmol, 1 equiv.) was dissolved in CH_2Cl_2 (2 ml) and treated with NaSCH_3 (29.2 mg, 0.416 mmol, 2 equiv.). The suspension was treated with concentrated HCl (4-5 drops) and stirred at RT. The reaction was complete after 30 min as determined by TLC. The reaction mixture was diluted with CH_2Cl_2 (40 ml), washed with sat. aq. NaHCO_3 (30

- 75 -

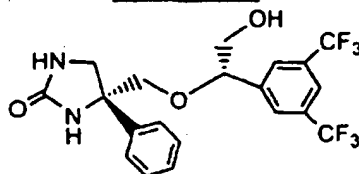
ml) and brine (30 ml), dried (Na_2SO_4), and concentrated. The crude product was purified by Biotage chromatography, eluting with 1:1 Hex/EtOAc to give the title compound (92.1 mg, 87%) as a 1:1 mixture of diastereomers. Electrospray MS $[\text{M}+1]^+$ 511.1

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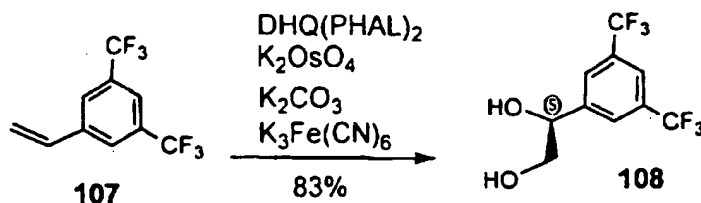
Example 97

A 2.3 diastereomeric mixture of Examples 94 and 95 (100 mg, 0.208 mmol, 1 equiv.) was dissolved in CH_3OH (3 ml), cooled to 0°C , and treated with concentrated HCl in CH_3OH (14 drops). This was warmed to RT and stirred to completion (24h).

- 10 The reaction mixture was diluted with CH_2Cl_2 (100 ml), washed with sat. aq. NaHCO_3 (30 ml) and then brine (30 ml), dried (Na_2SO_4), and concentrated. The crude mixture was purified by Biotage chromatography, eluting with 3:2 EtOAc/Hex (2% NEt_3) to give a product as a 3:2 mixture of diastereomers. Isolation by prep HPLC on a chiralcel OD column, eluting with 95:5 Hex/IPA gave the title compound ($t_R = 6.7$ min)
- 15 (29 mg, 28%). HRMS $[\text{M}+1]^+$ 495.1419

Example 98

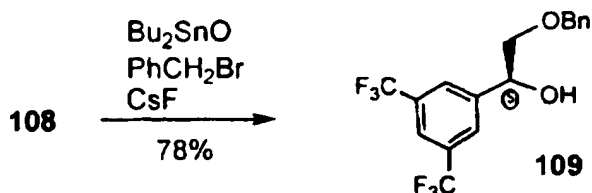
Step 1:



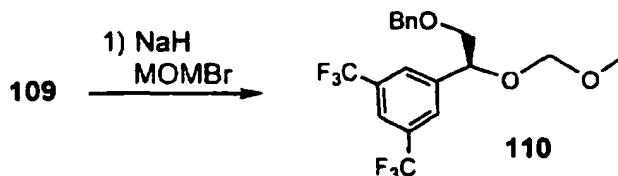
- 20 A solution of K_2CO_3 (31.0 g, 187.5 mmol, 3 equiv.), $\text{K}_3\text{Fe}(\text{CN})_6$ (61.75 g, 187.5 mmol, 3 equiv.), $(\text{DHQ})_2\text{PHAL}$ (2.44 g, 3.13 mmol, 0.05 equiv.), and $\text{K}_2\text{OsO}_4 \cdot 2\text{H}_2\text{O}$ (1.16 g, 3.13 mmol, 0.05 equiv.) in $t\text{BuOH}/\text{H}_2\text{O}$ 1:1 (750 ml) was stirred and cooled to 0°C . The suspension was treated with **107** (11.25 ml, 62.5 mmol, 1 equiv.) and stirred at 0°C for 3.5 h. The reaction mixture was treated with Na_2SO_3 (95 g, 750
- 25 mmol, 12 equiv.) and warmed to RT. The biphasic solution was separated and the aqueous layer was extracted with EtOAc (500 ml). Combined organic fractions were washed with brine (200 ml), dried (Na_2SO_4), and concentrated to a yellow solid. The

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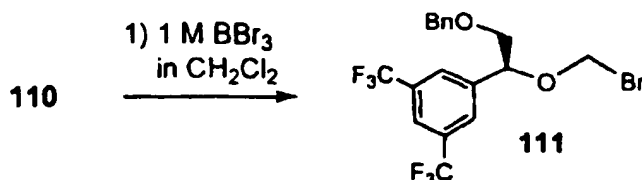
crude product was recrystallized from minimum 1,2-dichloroethane to give **108** (14.12 g, 83%) as a white solid. ^1H NMR (CD_3OD) δ 3.67 (d, J = 5.6 Hz, 2H), 4.85 (t, J = 5.6 Hz, 1H), 7.85 (s, 1H), 7.99 (s, 2H).

Step 2:

A mixture of **108** (4.0 g, 14.6 mmol, 1 equiv.) and dibutyltin oxide (3.64 g, 14.6 mmol, 1 equiv.) in toluene (60 ml) was refluxed for 2 h under a Dean Stark trap. The solution was concentrated to obtain a white solid. To this solid, CsF (4.30 g, 28.32 mmol, 1.94 equiv.) was added and the mixture was dried in vacuo overnight. Benzyl bromide (3.0 ml, 25.11 mmol, 1.72 equiv.) in dry DMF (60 ml) was added and the reaction mixture was stirred vigorously for 6 h. The solution was concentrated, taken up in EtOAc (200 ml) and the solid filtered off. The organics were extracted with water (4 x 100 ml), dried (Na_2SO_4), and concentrated. The crude product was purified by Biotage chromatography eluting with 9:1 Hex/EtOAc \rightarrow 4:1 Hex/EtOAc to give **109** (4.15 g, 78%) as a pale yellow oil. FAB MS $[M+1]^+$ 363.0.

Step 3:

NaH (834 mg, 20.85 mmol, 1.1 equiv.) was added to a stirred solution of **109** (6.90 g, 18.95 mmol, 1 equiv.) in THF (20 ml) under N_2 . The mixture was stirred for 1 h, then cooled to 0°C . MOMBr (2.63 g, 21.06 mmol, 1 equiv.) was added dropwise. The solution was warmed to RT and stirred for 1 h. The white mixture was filtered through a plug of celite and the filtrate was concentrated to give **110** (7.90 g, >95%) as a pure, yellow oil. ^1H NMR (CDCl_3) δ 3.36 (s, 3H), 4.55 (s, 2H), 4.61 (d, 7.0 Hz, 1H), 4.75 (d, 6.6 Hz, 1H), 4.92 (t, J = 4.8 Hz, 1H), 7.21-7.31 (m, 5H), 7.82 (s, 1H), 7.83 (s, 2H).

Step 4:

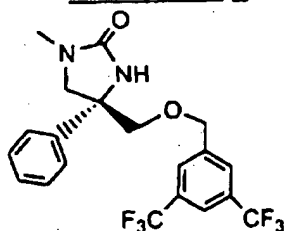
- 77 -

To an ice-cooled solution of **110** (5.85 g, 14.33 mmol, 3 equiv.) in CH_2Cl_2 (15 ml) was added a solution of 1 M BBr_3 in CH_2Cl_2 (1.64 ml, 1.64 mmol, 1 equiv.). The solution was warmed to RT and stirred for 4 h. The reaction mixture was concentrated to a crude brown oil **111** (6.1 g, 93%) which was used in the next step.

5 Step 5:

The title compound was prepared by a method analogous to Example 92 using **111** in place of **92** for step 4.

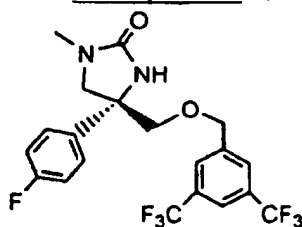
Example 99



- 10 The product of Example 92 (0.25g, 0.59 mmol, 1.0 equiv.) was taken up in 4 ml dry DMF and cooled to 0 °C in a ice bath. NaH (60% dispersion in mineral oil) (0.0179g, 0.59 mmol, 1.0 equiv.) was added to the reaction mixture, the solution was warmed to RT and stirred for 45 min. CH_3I (0.053 ml, 0.66 mmol, 1.1 equiv.) was added and reaction mixture was stirred at RT over night. The reaction was monitored
- 15 by TLC in 4/1 EtOAc/Hexane. The reaction did not go to completion, hence was quenched with H_2O (3x15ml). The mixture was extracted with EtOAc (2x15 ml) dried (Na_2SO_4), concentrated and dried on high vacuum. Purification using Biotage chromatography with 60/40 Hexane/EtOAc gave the title compound (0.127g, 50%). MS $[\text{M}+1]^+$ 433.1.

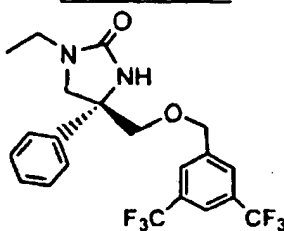
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Example 100



The title compound was prepared using a procedure similar to Example 99 using the product of Example 2B in place of Example 92. MS $[\text{M}+1]^+$ 451.1.

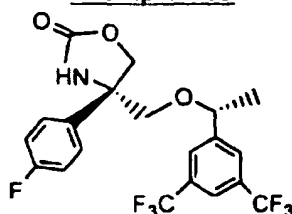
Example 101



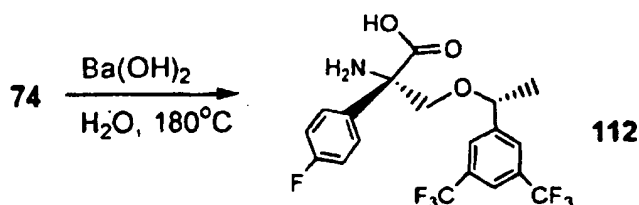
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The title compound was prepared using a procedure similar to Example 99 using ethyl iodide in place of CH_3I to give the title compound. MS $[\text{M}+1]^+$ of crude 447.1.

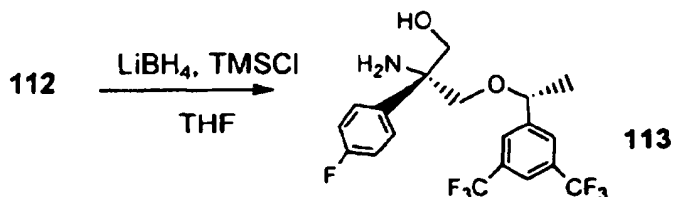
Example 102

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Step 1:

74 (6.0 g, 13.7 mmol, 1.0 equiv), $\text{Ba}(\text{OH})_2$ (20.8 g, 65.9 mmol, 4.8 equiv.) and H_2O (50 ml) were combined in a high pressure bomb equipped with a stirring bar. It was heated at 165 °C for 64 h, and then the temperature was increased to 180 °C. The bomb was heated for another 3 days, then cooled, and the crude product was treated with $\text{CH}_3\text{OH} (\text{NH}_3)$ to transfer the hardened crude to solution. The mass gradually dissolved using 10 (40-50 ml) portions with sonication each time. The resulting turbid solution was filtered and concentrated. The crude was then dissolved in ether, filtered and concentrated to give pale yellow solid 112 (6.2 g, >95%).

15

Step 2:

A solution of LiBH_4 (0.01 g, 0.46 mmol, 2.0 equiv) was treated with TMSCl (0.115 ml, 0.91 mmol, 4.0 equiv). After 5 min, 112 (0.1g, 0.227 mmol, 1.0 equiv.) was added as a powder and the mixture was stirred for 18 h. It was monitored by TLC 70:10:20 ($\text{EtOAc}:\text{Et}_3\text{N}:\text{CH}_3\text{OH}$). Upon completion, the reaction was quenched with 5ml CH_3OH and stirred for another 1 h. Purification by Biotage in 97:3 $\text{CH}_2\text{Cl}_2 / \text{CH}_3\text{OH} (\text{NH}_3)$ gave the product 113 (0.65 g, 67 %).

20

Step 3:

25

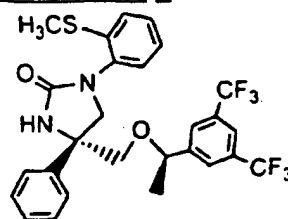
A solution of 113 (0.63 g, 0.148 mmol, 1.0 equiv) in dry CH_2Cl_2 (4ml) was treated with diisopropyl ethyl amine (0.077 ml, 0.444 mmol, 3.0 equiv.) followed by

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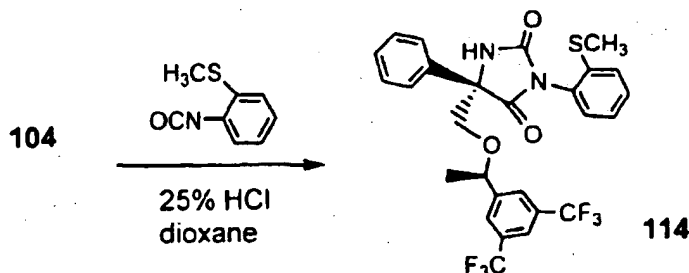
triphosgene (0.0176 g, 0.059 mmol, 0.4 equiv.). The reaction mixture was stirred at RT for 10 min. Monitoring by TLC 2/1 Hexane/EtOAc showed reaction completion. The crude product was purified using Biotage chromatography, eluting with 98/2 CH₂Cl₂ / CH₃OH to give the title compound as a white foam (0.58 g, 86 %). MS

5 [M+1]⁺ 452.

Example 103



Step 1



- 10 To a solution of **104** (0.225 g, 0.518 mmol) in CH₂Cl₂ (3 ml) was added 2-methylthiophenylisocyanate and the resulting solution was stirred for 12 h. The solvent was removed and the residue treated with dioxane/25% HCl (4 ml, 2:1, v/v) and heated at 90°C overnight. After cooling to RT, the solution was diluted with EtOAc and water and the layers separated. The aqueous layer was extracted with
- 15 EtOAc three times. The combined organic layers were washed with half saturated NaHCO₃ solution and brine, dried over MgSO₄, filtered and concentrated. Flash chromatography afforded **114** (0.265 g, 95%). Electrospray MS [M+1]⁺ 569.1

Step 2:

- 20 To solid AlCl₃ (346.7 mg, 2.6 mmol) at 0° C under N₂ was added LiAlH₄ (2.03 ml, 1M in Et₂O, 2.03 mmol). The resulting suspension was stirred for 10 min, then a solution of **114** (320 mg) in THF (19 ml) was cannulated into the hydride suspension dropwise. After 5 min, the solution was allowed to warm to 23 °C and stirred for 2 h. The reaction mixture was cooled to 0°C, quenched with 10 ml saturated sodium potassium tartrate, stirred at 23 °C for 2 h, then partitioned between water and EtOAc.
- 25 The organic layer was washed with brine, dried over MgSO₄, filtered and concentrated.

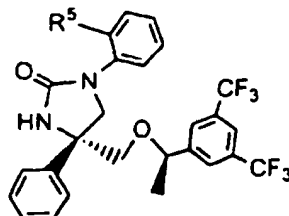
Step 3

- 80 -

A solution of the crude hydroxyurea from Step 2 in CH_2Cl_2 (3 ml) was treated with triethylsilane (1.5 ml) followed by addition of TFA (0.178 ml), stirred overnight, further treated with TFA (0.098 ml) and stirred for 4 h. The mixture was concentrated, then re-dissolved in CH_2Cl_2 and stirred with solid K_2CO_3 for 2 h. The solution was
 5 filtered and concentrated. Flash chromatography afforded the title urea. Electrospray MS $[\text{M}+1]^+$ 555.1.

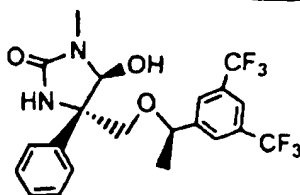
Examples 104-111

The following examples were prepared in a similar fashion to Example 103 using the appropriate isocyanate.

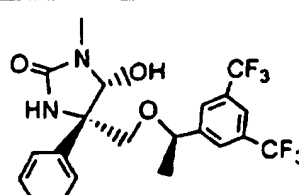


10

Example	R ⁵	MS $[\text{M}+1]^+$
104		539.1
105		545.1
106		527.1
107		539.1
108		537.1
109		539.1
110		569.1
111		528.1

Examples 112-113

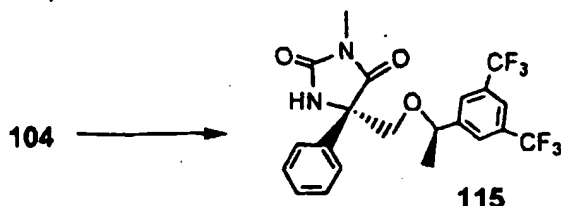
Ex. 112



Ex. 113

- 81 -

Step 1:



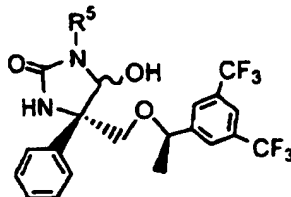
- 5 **104** (1.26 g, 2.9 mmol) was dissolved in CH_2Cl_2 (58 ml) at 0 °C, followed by addition of Et_3N (1.2 ml, 8.7 mmol). Triphosgene (0.35g, 1.13 mmol) was added in one portion. The solution was allowed to warm to 23 °C and stirred for 2 h. The reaction was then diluted with EtOAc, washed with 5% HCl(aq.) , half saturated NaHCO_3 solution and brine, then dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography to give **115** (1.31g, 98%).

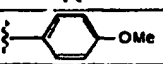
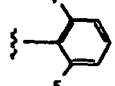
10 Step 2:

Using a procedure similar to Example 94, step 2, a mixture of Examples 112 and 113 was prepared. Separation by silica gel chromatography using gradient of hexane to EtOAc/ NEt_3 9:1 provided the title compounds.

Examples 114-115

- 15 The following examples were prepared in a similar fashion to Example 103, steps 1-2, omitting step 3 and using the appropriate isocyanate:

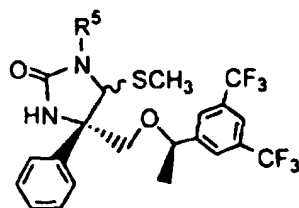


Example	R^5	MS $[\text{M}+1]^+$
114		555.1
115		561.1

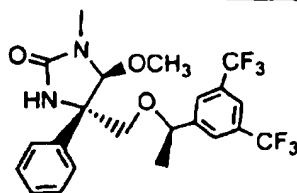
Examples 116-117

- 20 The following examples were prepared in a similar fashion to Example 96, using 116 to prepare Example 116 and Example 115 to prepare Example 117 in place of hydroxyureas from Examples 94 and 95:

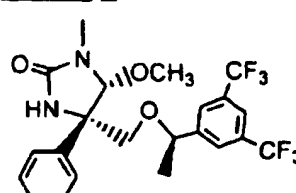
- 82 -



Example	R ⁵	MS [M+1] ⁺
116	Me	493.1
117		591.1

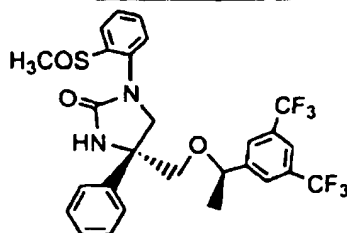
Examples 118-119

Ex. 118



Ex. 119

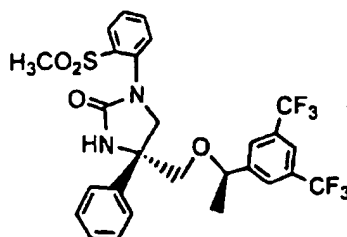
To the alcohol mixture of Examples 112 and 113 was added saturated dry HCl in CH₃OH (5 ml, precooled to -20 ° C). The solution was stirred at 23 °C for 2 h, then poured into 10% Na₂CO₃ (25 ml). The aqueous layer was extracted with EtOAc (3 X 30 ml). The combined organic layer was washed with brine, dried over Na₂SO₄, filtered and concentrated under reduced pressure. Flash chromatography afforded an inseparable mixture of two methyl ethers. The mixture was then separated on HPLC using Chiralpak OD eluted with (95:5) Hexane/iPA. Electrospray MS [M+1]⁺ 477.1

Example 120

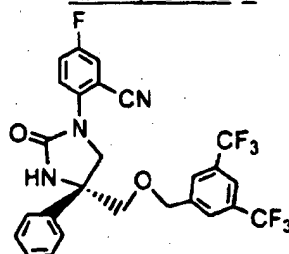
To a solution of Example 103 (170 mg, 0.306 mmol) in CH₂Cl₂ was added solid Oxone (1.14g, 17.4 mmol), the resulting suspension was stirred vigorously for 20 h, then quenched with saturated NaHCO₃ solution and extracted with EtOAc. The organic layers were washed with brine, dried over Na₂SO₄ and concentrated. Flash chromatography afforded the title compound. Electrospray MS [M+1]⁺ 571.1

Example 121

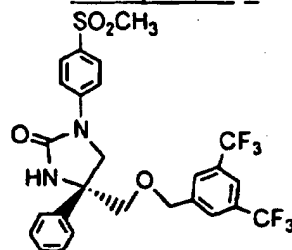
- 83 -



To a solution of Example 120 in CH₂Cl₂ was added *m*-CPBA and the resulting solution was stirred for 30 h, then quenched with NaHCO₃ solution and extracted with EtOAc. The organic layers were washed with brine, dried over Na₂SO₄ and concentrated. Flash chromatography afforded the title compound. Electrospray MS [M+1]⁺ 587.1.

Example 122

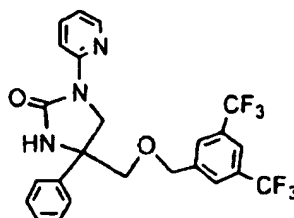
To Example 92 (0.150 g, 0.36 mmol), CsF (0.109 g, 0.72 mmol) in DMF (3.5 ml) at 0°C and NaH 0.018 g (60% in mineral oil) was added. After stirring for 10 min, 2,5-difluorobenzonitrile (0.0587 g, 0.0378 mmol) was added in small portions. The solution was allowed to warm to 23 °C and was stirred overnight. The reaction was quenched with saturated NH₄Cl solution, extracted with EtOAc, and the combined organic layer was washed with water and brine, dried over Na₂SO₄ and concentrated. Flash chromatography afforded the title compound (0.051g, 26%). Electrospray MS [M+1]⁺ 538.1.

Example 123

In a procedure similar to Example 122, 4-F-phenylmethylsulfone was used in place of 2,5-difluorobenzonitrile to obtain the title compound. Electrospray MS [M+1]⁺ 573.1.

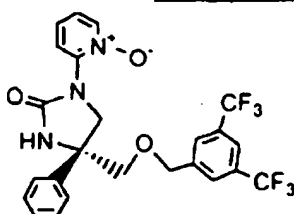
Example 124

- 84 -

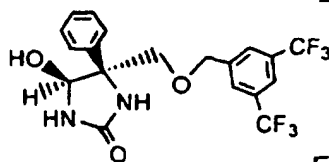


In a procedure similar to Example 122, 2-F-pyridine was used in place of 2,5-difluorobenzonitrile and Example 1 was used in place of Example 92 to obtain the title compound. Electrospray MS $[M+1]^+$ 496.1

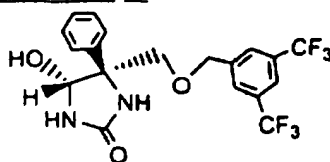
5

Example 125

In a procedure similar to Example 122, 2-F-pyridine was used in place of 2,5-difluorobenzonitrile in step 1, followed by a procedure similar to that used in Example 121 for the oxidation to the N-oxide to obtain the title compound. Electrospray MS

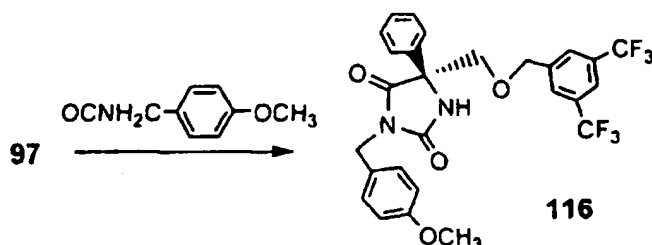
10 $[M+1]^+$ 512.1.Examples 126-127

Ex. 126



Ex 127

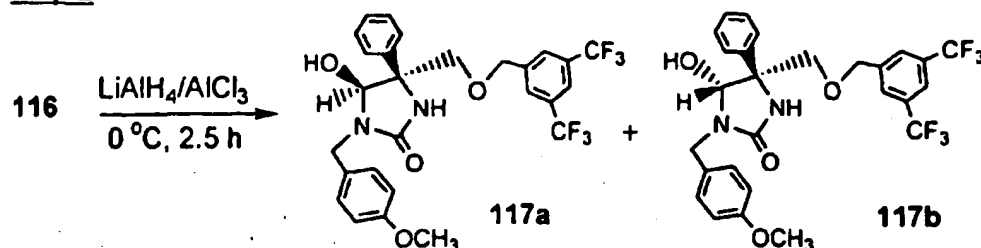
Step 1:



15 A mixture of hydantoin (prepared from 97 using procedures similar to Example 61, step 6) (5.0 g, 11.6 mmol, 1 equiv.), and *p*-methoxybenzylisocyanate (2.5 ml, 17.4 mmol, 1.5 equiv.) in dry dioxane (20 ml) was stirred at RT for 3 h, then 3N aqueous HCl (20 ml) was added and the mixture stirred at 90 °C for 14 h. The mixture was poured into 250 ml EtOAc and washed with H₂O (2 x 125 ml). The organic layer was

20 dried over anhydrous Na₂SO₄, filtered and concentrated to give 6.5 g of the crude racemic 116.

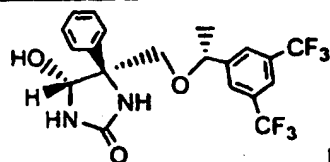
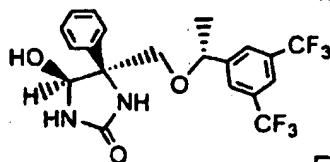
- 85 -

Step 2

- LiAlH₄ (35 ml of 1 M solution in ether, 35 mmol, 3 equiv.) was added slowly to AlCl₃ (6.3 g, 47.06 mmol, 4 equiv.) at 0°C, and stirred for 10 min, then a solution of 116 (6.5 g, 11.76 mmol, 1 equiv.) in dry THF (70 ml) was carefully added. After stirring at 23 °C for 2.5 h, the mixture was cooled to 0°C, quenched slowly with 30 ml saturated aqueous sodium potassium tartarate and stirred for 14 h at 23 °C. The mixture was diluted with water (100 ml), and extracted with EtOAc (2 x 200 ml). The organic layers were dried over anhydrous Na₂SO₄, filtered and concentrated.
- Purification using 400 ml silica and eluting with 2/1 hexane/EtOAc provided 4.62 g of the product as white solid.

Step 3:

- To a suspension of 117a/b (4.62 g, 8.33 mmol, 1 equiv.) in CH₃CN/water (150 ml, 2:1) at RT was added ceric ammonium nitrate (18.27 g, 33.33 mmol, 4 equiv.). After stirring at 23 °C for 1 h, the mixture was poured into 300 ml EtOAc/ 150 ml saturated aq. NaHCO₃ and filtered through a frit. The organic layer was isolated, the aqueous layer was washed with EtOAc (1 x 300 ml) and the combined organic layers were dried over anhydrous Na₂SO₄, filtered and concentrated. Purification using the Biotage silica gel system, eluting with 1/1 hexane/EtOAc, 1 L, followed by 5% CH₃OH/EtOAc, 1 L, provided 2.0 g of the product as a mixture of two isomers. HPLC separation on chiralpak AD column, eluting with (90/10) hexane/IPA mixture gave Example 127 FAB, (M⁺) = 435.0, and Example 126, FAB, (M⁺) = 435.0.

Examples 128-129Step 1:

- A mixture of 105 (0.976 g, 2.19 mmol, 1 equiv.), K₂CO₃ (0.453 g, 3.28 mmol, 1.5 equiv.) in dry DMF (10 ml) was stirred at RT for 30 min. *p*-Methoxy-benzyl chloride (0.34 ml, 2.5 mmol, 1.15 equiv.) was added at once and the resulting mixture was stirred at 23°C for 14 h. The mixture was then poured into EtOAc (150 ml) and washed with H₂O (3 x 100 ml) and saturated aq. NaCl (100 ml). The organic layer

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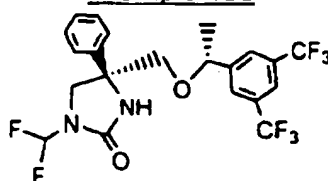
was dried over Na_2SO_4 , filtered and concentrated to give 1.27 g of the crude product. It was used in the next step without further purification.

Step 2:

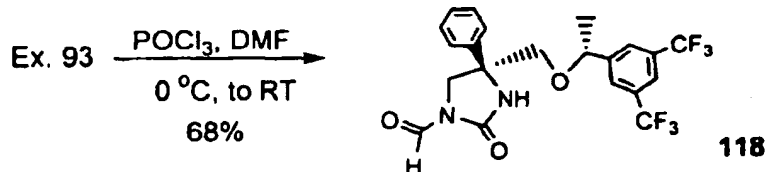
- The product of step 1 was converted to Examples 128 and 129 using procedures similar to Examples 126 and 127, steps 2-3, using LAH in place of LAH/ AlCl_3 in step 2 to provide 0.14 g of the products, obtained as mixture of two isomers. HPLC separation on chiralpak AD column using (85/15) hexane/IPA mixture gave 80 mg (29% yield) of Example 129, FAB, (M^{+1}) 449.2 and 30 mg (11% yield) of Example 128 FAB, (M^{+1}) 449.2.

10

Example 130



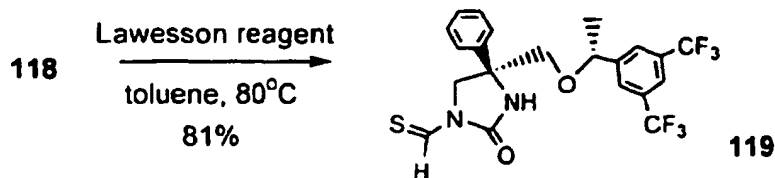
Step 1:



- To a solution of Example 93 (0.1 g, 0.23 mmol, 1.0 equiv.) in dry DMF (0.5 ml) cooled to 0 °C, POCl_3 (0.024 ml, 0.254 mmol, 1.1 equiv.) was added slowly. The mixture was warmed to RT, stirred for 30 min, and poured into 5 g ice. The resultant mixture was poured into water (100 ml) and extracted with EtOAc (100 ml). The organic layer was separated, washed with saturated aq. NaCl (1 x 100 ml), dried over anhydrous Na_2SO_4 and concentrated. Flash chromatography over 200 ml silica using (1) 4/1 hex/EtOAc, and (2) 1/1 hex/EtOAc gave 0.070 g (66% yield) of **118** as solid. MS (M^{+1}) 461.1.

20

Step 2:



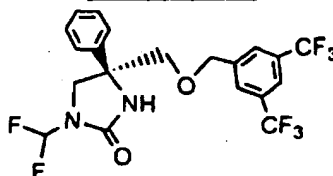
- 118** (0.1 g, 0.22 mmol, 1.0 equiv.) and Lawesson's reagent (0.044 g, 0.108 mmol, 0.5 equiv.) in toluene (1 ml) were heated at 80 °C for 0.5 h. The solvent was evaporated and the residue purified by Biotage chromatography using 15% EtOAc/hexane to obtain **119** as a yellow foam, 0.085g (81% yield).

25

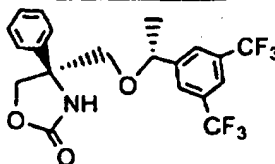
Step 3:

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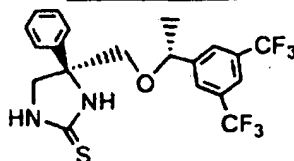
- 119 (0.3 g, 0.626 mmol, 1.0 equiv.) was taken up in CH_2Cl_2 (3 ml) at RT. DAST (0.17 ml, 1.25 mmol, 2.0 equiv.) was added slowly and the mixture was stirred at RT overnight. The reaction was slowly quenched with saturated aq. NaHCO_3 (5 ml), the mixture was poured into saturated aq. NaHCO_3 (100 ml) and CH_2Cl_2 (100 ml) added. The organic layer was separated, dried over anhydrous Na_2SO_4 , filtered and concentrated. The residue was purified by Biotage chromatography using 4/1 hex/EtOAc to obtain the title compound, 0.080 g (27% yield). MS (M^+) = 483.1.1.

Example 131

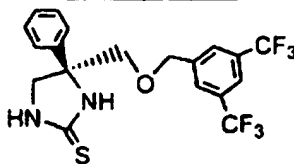
- 10 Using procedures similar to Example 130, substituting Example 92 for Example 93, the title compound was prepared. Electrospray MS [$M+1$] $^+$ 469.1

Example 132

- 15 Using procedures similar to those in Example 102, and substituting 97 for 74 in step 1, the title compound was prepared. Electrospray MS [$M+1$] $^+$ 434.1.

Example 133

- 20 Example 93 (0.633 g, 1.465 mmol, 1.0 equiv.), Lawesson's reagent (0.81 g, 2.04 mmol, 1.37 equiv.) and toluene (12 ml) were heated at 85 °C for 1.5 h, then cooled to RT and concentrated. The residue was purified by silica gel chromatography using 15% EtOAc/hex, then 10% EtOAc/ CH_2Cl_2 to give 0.61 g (93% Yield) of the title compound. MS (M^+) = 449.1.

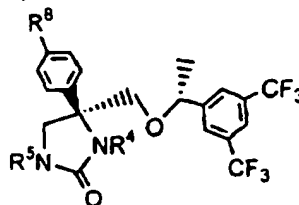
Example 134

- 25 A procedure similar to that used in Example 133, using Example 92 in place of Example 93, provided the title compound. MS (M^+) = 435.1

- 88 -

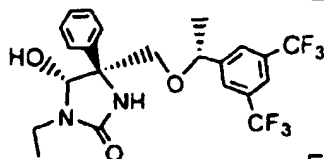
Examples 135-139

The compounds were prepared using procedures similar to those used in Example 99, using Example 92 and the appropriate alkyl iodide. For Example 139, Example 76 was used as the starting cyclic urea.

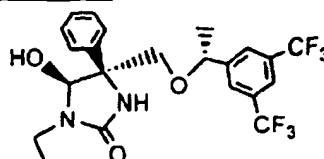


5

Example	R ⁵	R ⁴	R ⁸	MS [M+1] ⁺
135	Me	H	H	447.1.
136	Et	H	H	461.1.
137	H	Et	H	461.1.
138	<i>i</i> Pr	H	H	475.1.
139	Et	H	F	479.1.

Examples 140-141

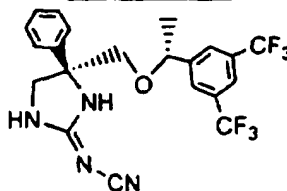
Ex. 140



Ex. 141

The title compounds were prepared using procedures similar to those in Example 94, using ethyl isocyanate in place of methyl isocyanate. Purification via HPLC (Chiralpak AD column using 98/2 hex/isopropanol) provided Example 140 MS [M+1]⁺ 477.1 and Example 141 MS [M+1]⁺ 477.1.

10

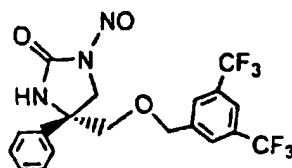
Example 142

Example 133 (0.61 g, 1.36 mmol, 1.0 equiv.) in THF (20 ml) was treated with CH₃I (0.10 ml, 1.63 mmol, 1.2 equiv.), stirred for 14 h, then concentrated. The crude was dissolved in CH₃OH (20 ml), treated with NH₂CN (0.37 g, 8.84 mmol, 6.7 equiv.) and heated to 60 °C for 14 h. The mixture was concentrated and purified by silica gel chromatography, using 1/1 EtOAc/hex to give the title compound, 0.1 g (17% yield) as a white solid. MS (M⁺) = 457.1.

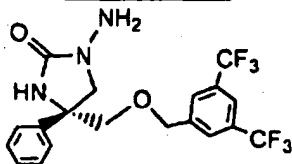
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Example 143

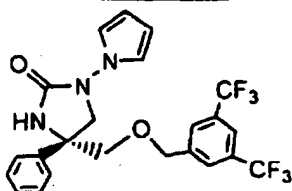
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- To Example 92 (2.0 g, 4.78 mmol) in THF (15 ml) at 0° C was added NaNO₂ (0.39 g, 5.7 mmol) in H₂O (7.5 ml), then H₂SO₄ (conc.) (1 ml) was added slowly. The solution was allowed to warm to 23°C and stirred for 1 h. It was then diluted with water and extracted with EtOAc. The organic layers were washed with NaHCO₃ solution, brine, dried over Na₂SO₄ and concentrated to give the title compound. Electropray MS [M+1]⁺ 448.1.

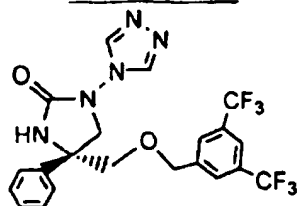
Example 144

- To Example 143 (0.95 g, 2.1 mmol) in Et₂O (10 ml) at 0° C was added dropwise LiAlH₄ (4.2 ml, 1M in Et₂O). The solution was allowed to warm to 23°C and stirred for 2 hr. It was quenched with saturated K₂Na tartrate solution and then partitioned between water and EtOAc. The aqueous layer was extracted with EtOAc. The combined organic layers were washed with brine, dried over Na₂SO₄ and concentrated to give the title compound. Electropray MS [M+1]⁺ 434.1

Example 145

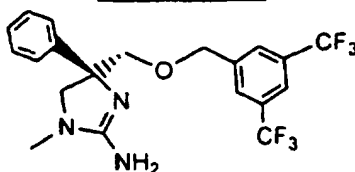
- To Example 144 (0.3 g, 0.69 mmol) in acetic acid (3 ml) was added 2,5-dimethoxy-3-tetrahydrofuran (3 ml) and the mixture was heated at 70° C for 1.5 h. It was cooled to 23°C and diluted with EtOAc. The organic layers were washed with NaHCO₃, brine, dried over Na₂SO₄ and concentrated. A stream of N₂ was introduced into the residue to remove excess 2,5-dimethoxy-3-tetrahydrofuran. The crude product was purified by silica gel chromatography to give the title compound. Electropray MS [M+1]⁺ 484.1.

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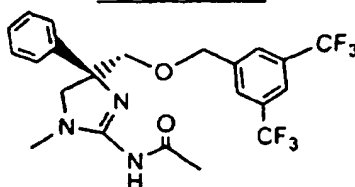
Example 146

Example 144 (0.65 g, 1.5 mmol) in pyridine (15 ml) was concentrated under reduced pressure at 50° C using a water bath. The procedure was repeated twice.

- 5 The residue was treated with 1,2-diformylhydrazine (0.34g, 3.6 mmol), followed by the addition of pyridine (7.5 ml), Et₃N (1.5 ml) and TMSCl (3 ml). The thick paste was heated at 80 °C under N₂ for 65 h, then concentrated under reduced pressure to give a yellow residue. The residue was partitioned between water and EtOAc. The aqueous layer was extracted with EtOAc. The combined organic layers were washed
- 10 with brine, dried over Na₂SO₄ and concentrated to give a crude product. Further purification using biotage followed by a prep TLC afforded the title compound. Electrospray MS [M+1]⁺ 486.1.

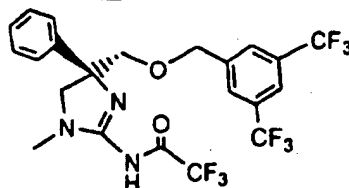
Example 147

- 15 The title compound was prepared in a method analogous to Example 73, using 89 in place of 6. The title compound was obtained in 95% yield. Electrospray MS [M+1]⁺ 432.1.

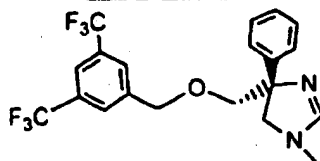
Example 148

- 20 To a cooled solution of Example 147 (0.3 g, 0.70 mmol), DEC (0.14 g, 0.73 mmol), HOBT (0.1 g, 0.74 mmol) and NaSO₄ (0.59 g) in anhydrous CH₂Cl₂ (4.5 ml), glacial acetic acid (0.05 ml, 0.87 mmol) was added followed by Et₃N (0.1 ml, 0.72 mmol). The reaction was allowed to react for 18 h at RT, then quenched with brine. The aqueous solution was extracted with CH₂Cl₂ (50 ml X 3). The combined organic
- 25 layer was dried, filtered and concentrated. The pure title compound was obtained through flash chromatography, eluting with 2% (1:9) NH₄OH-CH₃OH in CH₂Cl₂ to give the final product in 39% yield. Electrospray MS [M+1]⁺ 474.1.

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Example 149

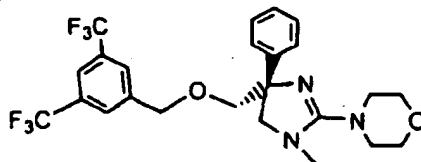
To a cooled solution of Example 147 (0.2 g, 0.46 mmol) in anhydrous CH_2Cl_2 (5 ml) at 0 °C, trifluoroacetic anhydride (0.09 ml, 0.64 mmol) neat was added followed by Et_3N (0.08 ml, 0.57 mmol). The reaction was allowed to react for 18 h at RT. Volatile solvents were evaporated. The pure title compound was obtained through flash chromatography, eluting with 50% EtOAc in hexane to give the final product in 33% yield. Electrospray MS $[\text{M}+1]^+$ 528.1.

Example 150

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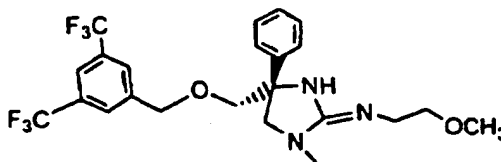
A mixture of 89 (0.38 g, 0.93 mmol) and N, N-dimethylformamide dimethyl acetal (0.11g, 0.93 mmol) was heated to 60 °C for 18 h. The reaction mixture was purified by chromatography, eluting with 3.5 % NH_3 - CH_3OH (1:9)/96.5 % CH_2Cl_2 to give the title compound later as a HCl salt (0.2 g, 50 %). FAB MS $[\text{M}+1]$ 417.

15

Example 151

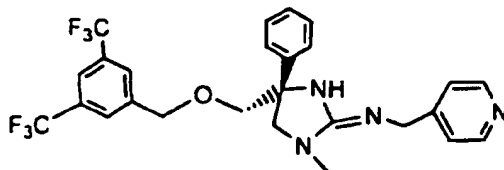
Using a method analogous to Example 82, using morpholine in place of 7 M NH_3 in CH_3OH , the title compound was obtained as a solid in a 36 % yield. FAB MS $[\text{M}+1]$ 502.

20

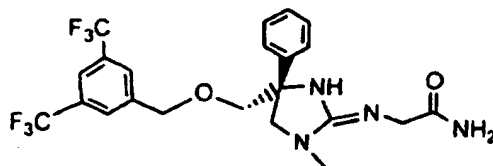
Example 152

Using a method analogous to that Example 82, using 2-methoxyethyl-amine in place of 7 M NH_3 in CH_3OH , the title compound was obtained as a solid in a 80 % yield. FAB MS $[\text{M}+1]$ 490.1.

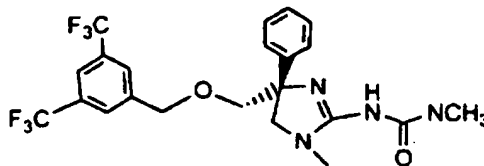
- 92 -

Example 153

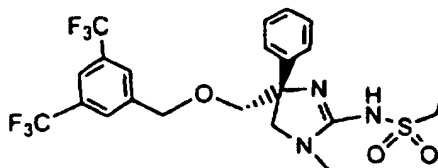
Using a method analogous to Example 82, using 4-(aminomethyl)pyridine in place of 7 M NH_3 in CH_3OH , the title compound was obtained as a solid in a 26 % yield. FAB MS $[\text{M}+1]$ 523.4.

Example 154

Using a method analogous to Example 82, using glycine in place of 7 M NH_3 in CH_3OH , the title compound was obtained as a solid in a 14 % yield. FAB MS $[\text{M}+1]$ 489.2.

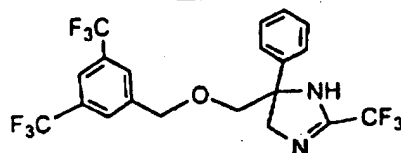
Example 155

A solution of Example 82 (0.2 g, 0.464 mmol) in CH_2Cl_2 (2 ml) and isocyanate (53 mg, 0.93 mmol) was stirred under N_2 at RT for 18 h. After work-up, a solid was obtained as a crude product which was then purified by chromatography, eluting with 2 % NH_3 - CH_3OH (1:9) / 98 % CH_2Cl_2 to give the title compound as a solid (55 mg, 24 %). FAB MS $[\text{M}+1]$ 489.3.

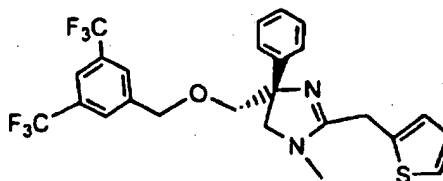
Example 156

Using a method analogous to Example 155, using ethane sulfonyl chloride in place of isocyanate, the title compound was obtained later as a HCl salt in a 25 % yield. FAB MS $[\text{M}+1]$ 524.3.

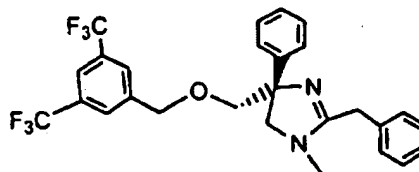
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Example 157

Using a method analogous to Example 87, using **6** in place of **89** and TFA in place of acetic acid, the title compound was later obtained as a HCl salt in a 34 %
5 yield. FAB MS [M+1] 471.

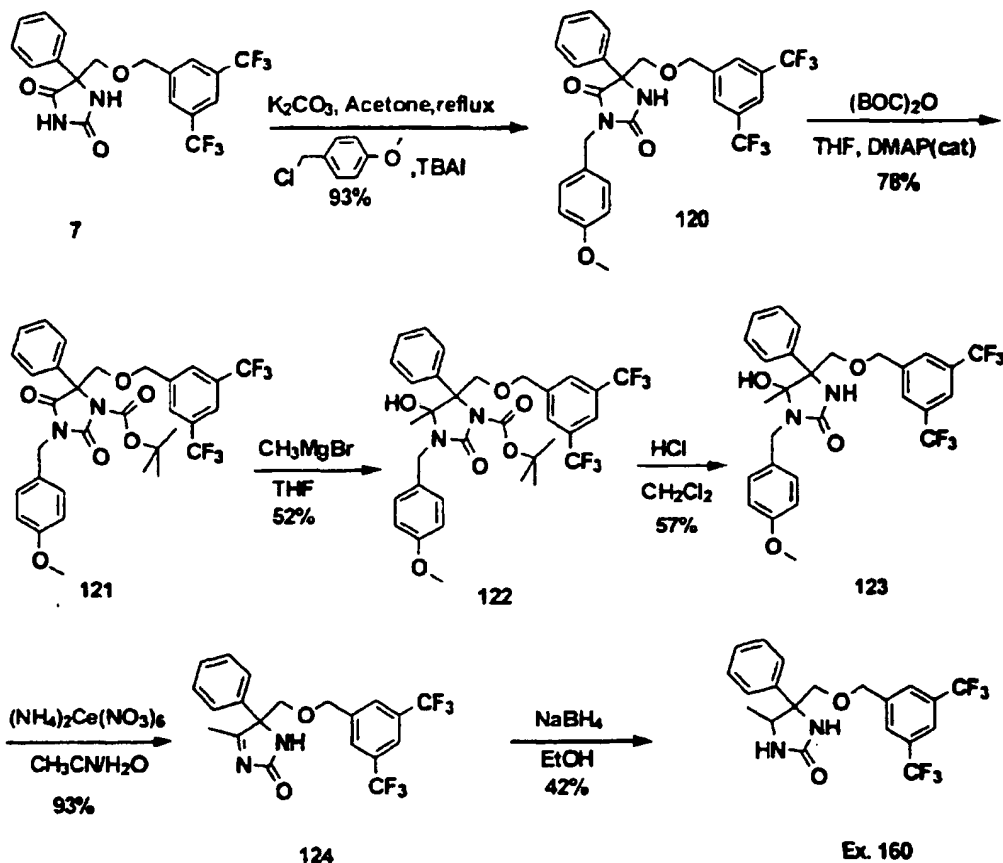
Example 158

Using a method analogous to Example 89, using 2-thiophene acetic acid in place of acetic acid, the title compound was later obtained as a HCl salt in a 24 %
10 yield. FAB MS [M+1] 512.9.

Example 159

Using a method analogous to Example 89, using phenyl acetic acid in place of acetic acid, the title compound was later obtained as a HCl salt in a 15 % yield. FAB
15 MS [M+1] 507.1.

Example 160



5

Step 1 A mixture of hydantoin 7 (9.2 g, 21.3 mmol, 1 equiv.), K_2CO_3 (8.8g, 63.9 mmol, 3.0 equiv.) and tetrabutylammonium iodide (0.79 g, 2.13 mmol, 0.1 equiv.) in 180 mL acetone was stirred at room temperature for 30 minutes, treated with p -methoxybenzyl chloride (3.25 mL, 23.9 mmol, 1.12 equiv.) at reflux for 2 hours, then cooled to room temperature and filtered through sintered glass funnel. The filtrate was concentrated to give pale yellow solid, which was washed with small amount of ice-cold Et_2O (2 X 5 mL) to give 11 g of the crude racemic p -methoxybenzyl (pmb)-hydantoin 120. It was used in the next step without further purification.

10

Step 2 A mixture of pmb-hydantoin 120 (6.0 g, 10.9 mmol, 1 equiv.), 4-(dimethylamino)pyridine (DMAP) (0.02 g, 0.16 mmol, 1.5% equiv.) in 60 mL dry THF was stirred at room temperature for 1.5 hours. The solvent was evaporated under vacuum. The residue was taken up in CH_2Cl_2 (150 mL) and washed with saturated sodium bicarbonate (50 mL). The organic layer was dried

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over anhydrous Na_2SO_4 , then concentrated. The crude was purified with Biotage (EtOAc/Hexane=10%) to give 5.5 g (77.7% yield) of BOC-pmb-hydantoin **121** as white solid.

- Step 3** A solution of BOC-pmb-hydantoin **121** (540 mg, 0.83 mmol, 1 equiv.) in 10 mL dry THF was treated with 3.0 M methylmagnesium bromide Et_2O solution (0.42 mL, 1.25 mmol, 1.5 equiv.) at 0 C. After addition, the reaction was warmed up to room temperature gradually and stirred for another hour. THF was evaporated under vacuum. The residue was taken up with 100 mL CH_2Cl_2 , then washed with 10 mL of saturated aq. NaHCO_3 solution. The resulting white solid was filtered off. The filtrate was concentrated and purified by Biotage (EtOAc/Hexane=20%) to give 290 mg (52.3% yield) of Methyl-hydroxy-BOC-pmb-urea **122** as white solid.

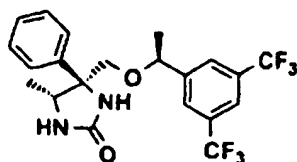
- Step 4** A solution of methyl-hydroxy-BOC-pmb-urea **122** (310 mg, 0.46 mmol, 1 equiv.) in 5 mL of dry CH_2Cl_2 was treated with 1.16 mL of 4.0 M HCl in 1,4-dioxane at 0 C. The reaction was allowed to warm up to room temperature and stirred overnight. The crude product was taken up with Et_2O , then washed with 3 mL of saturated NaHCO_3 . The aqueous layer was further extracted with ether. The combined organic layer was dried over Na_2SO_4 , then concentrated, and purified with Biotage (EtOAc/Hexane=15%) to give 150 mg (56.5% yield) of Methyl-hydroxy-pmb-urea **123** as white solid.

- Step 5** A white suspension of Methyl-hydroxy-pmb-urea **123** (1g, 1.76 mmol, 1 equiv.) in 22.5 mL of CH_3CN and 6.75 mL of water was treated with ceric ammonium nitrate (7.72 g, 14.0 mmol, 8 equiv.). Stirred at room temperature for 8 hours, then partitioned between 300 mL of EtOAc and 100 mL of saturated aq. NaHCO_3 . The yellow solid was filtered off and aqueous layer was further extracted with 2 x 100 mL EtOAc. Combined organic layers were dried over anhy. Na_2SO_4 , filtered and concentrated. Flashed over Biotage (EtOAc/Hexane/ Et_3N =1:1:2%) to give 703 mg (93% yield) of **124** as white solid.

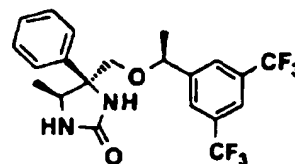
- Step 6** A solution of **124** (312 mg, .93 mmol, 1 equiv.) in 8 mL EtOH was reacted with NaBH_4 (600 mg, 15.9 mmol, 17 equiv.) at room temperature for 2 days. The crude was partitioned between 2 X 100 mL of CH_2Cl_2 and 100 mL saturated aq. NaHCO_3 , then washed with 80 mL of Brine. The combined organic layer was dried over anhy. Na_2SO_4 , filtered, concentrated to give 300 mg of liquid as crude. It was purified with Biotage (EtOAc/Hexane=30%) to give two diastereomers (100mg and 70 mg each), which were separated on HPLC with a ChiralCel OD column to give the 4 stereoisomers of example **160**. MS: (M+1)= 433.

Example 161

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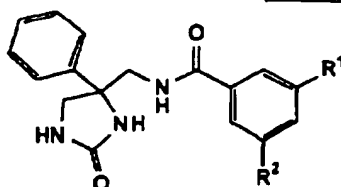
Example 161a



Example 161b

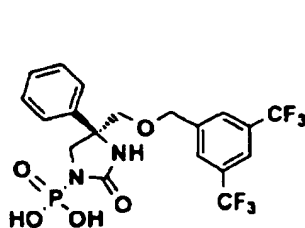
Examples 161a and 161b were synthesized using procedure similar to example 160 starting from optical pure hydantoin 105. MS: (M+1)=447.

5

Examples 162-164

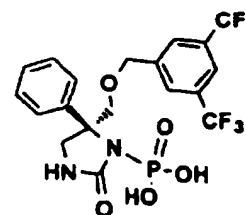
Example	R ¹	R ²	MS [M+1] ⁺
162	F	CF ₃	382.1.
163	Cl	Cl	364.1.
164	CF ₃	CF ₃	432.1

The title compounds were prepared using the acylation procedure similar to those used in example 21 using amine 39 and the appropriate substituted benzoyl chloride to provide the title benzamides.

Example 165

• 2 N-methylglucamine

Example 165a



• 2 N-methylglucamine

Example 165b

15

Step 1

A solution of Example 92 (224 mg, 0.536 mmol, 1 equiv.) in THF (12 mL) at 0°C was treated with 2.5 M n-BuLi (215 µL, 0.536 mmol, 1 equiv.). The resulting mixture was stirred cold for 5 min. Tetrabutyl pyrophosphate (405 mg, 0.751 mmol, 1.4 equiv.) was added to the reaction mixture as a solid in one portion. The

20

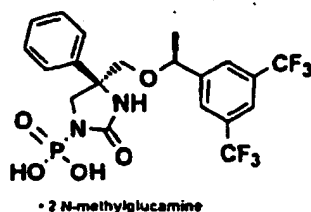
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cooling bath was removed and the reaction was stirred at room temperature for 45 minutes at which point a white suspension formed. The reaction was quenched with sat. aq. NaHCO_3 (20 mL) and extracted with EtOAc (2 x 30 mL). The solution was dried (Na_2SO_4), filtered, and concentrated to give crude product (461 mg). This was not purified and used crude in the next step.

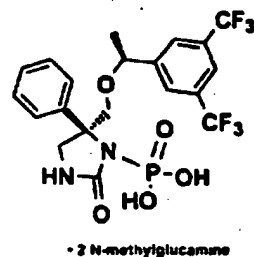
Step 2

A solution of the product of step 1 (364 mg, 0.536 mmol, 1 equiv.) in MeOH (10 mL), a solution of N-Me-D-glucamine (206 mg, 1.072 mmol, 2 equiv.) in H_2O (2 mL), and 10% Pd/C (29 mg) were combined and the mixture was hydrogenated at 40 psi for 2 h. The reaction mixture was filtered through a pad of Celite and rinsed with MeOH (80 mL). The solution was concentrated under reduced pressure and the crude product redissolved in MeOH (5 mL). PrOH (25 mL) was added to the solution and the resulting mixture was aged at room temperature for 30 min to form a white precipitate. The precipitate was filtered, washed with PrOH (15 mL) and EtOAc (15 mL), and dried. The solid was partitioned between EtOAc (30 mL) and H_2O (30 mL) and an emulsion formed. The emulsion was transferred in 10-12 mL portions to centrifuge tubes and centrifuged at 3000 rpm for 15 min. Decanted away the organic layer. The aqueous layers were combined, filtered, and lyophilized to provide example 165 (306 mg, 64% yield, 1:1 mixture of regioisomers 165a and 165b).

Example 166



Example 166a



Example 166b

Using procedures similar to those in example 165 and substituting Example 93 for Example 92, the title compounds were prepared.

Compounds of formula I have been found to be antagonists of the NK_1 receptor and of the effect of Substance P at its receptor site, and are therefore useful in treating conditions caused or aggravated by the activity of said receptor.

The *in vitro* and *in vivo* activity of the compounds of formula I can be determined by various procedures known in the art, such as a test for their ability to inhibit the activity of the NK_1 agonist Substance P. % Inhibition of neurokinin agonist activity is the difference between the percent of maximum specific binding (MSB) and

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100%. The percent of MSB is defined by the following equation, wherein "dpm" is disintegrations per minute:

$$\% \text{ MSB} = \frac{(\text{dpm of unknown}) - (\text{dpm of nonspecific binding})}{(\text{dpm of total binding}) - (\text{dpm of nonspecific binding})} \times 100$$

5 The concentration at which the compound produces 50% inhibition of binding is then used to determine the inhibition constant (k_i) using the Chang-Prusoff equation.

In addition, functional antagonism of calcium channel activity is measured using FLIPR technology known to those skilled in the art.

10 *In vivo* activity is measured by inhibition of agonist-induced foot tapping in the gerbil, as described in Science, (1998), 281, p. 1640-1695.

It will be recognized that compounds of formula I exhibit NK_1 antagonist activity to varying degrees, e.g., certain compounds have strong NK_1 antagonist activity, while others are weaker NK_1 antagonists.

15 Compounds of the present invention exhibit a range of activity: K_i values range from about 0.1 to 1000 nM, with K_i values of about 0.1 to 100 being preferred and K_i values of 0.1 to 25 nM being more preferred. Most preferred are compounds having a $K_i \leq 10$ nM for the NK_1 receptor.

20 For preparing pharmaceutical compositions from the compounds described by this invention, inert, pharmaceutically acceptable carriers can be either solid or liquid. Solid form preparations include powders, tablets, dispersible granules, capsules, cachets and suppositories. The powders and tablets may be comprised of from about 5 to about 95 percent active ingredient. Suitable solid carriers are known in the art, e.g. magnesium carbonate, magnesium stearate, talc, sugar or lactose. Tablets, powders, cachets and capsules can be used as solid dosage forms suitable for oral
25 administration. Examples of pharmaceutically acceptable carriers and methods of manufacture for various compositions may be found in A. Gennaro (ed.), Remington's Pharmaceutical Sciences, 18th Edition, (1990), Mack Publishing Co., Easton, PA.

30 Liquid form preparations include solutions, suspensions and emulsions. As an example may be mentioned water or water-propylene glycol solutions for parenteral injection or addition of sweeteners and opacifiers for oral solutions, suspensions and emulsions. Liquid form preparations may also include solutions for intranasal administration.

35 Aerosol preparations suitable for inhalation may include solutions and solids in powder form, which may be in combination with a pharmaceutically acceptable carrier, such as an inert compressed gas, e.g. nitrogen.

Also included are solid form preparations which are intended to be converted, shortly before use, to liquid form preparations for either oral or parenteral administration. Such liquid forms include solutions, suspensions and emulsions

The compounds of the invention may also be deliverable transdermally. The
5 transdermal compositions can take the form of creams, lotions, aerosols and/or emulsions and can be included in a transdermal patch of the matrix or reservoir type as are conventional in the art for this purpose.

Preferably the compound is administered orally.

Preferably, the pharmaceutical preparation is in a unit dosage form. In such
10 form, the preparation is subdivided into suitably sized unit doses containing appropriate quantities of the active component, e.g., an effective amount to achieve the desired purpose.

The quantity of NK₁ receptor antagonists in combination with selective serotonin reuptake inhibitors in a unit dose of preparation may be varied or adjusted
15 from about 10 mg to about 300 mg of NK₁ receptor antagonists with about 10 mg to about 100 mg of SSRI. A further quantity of NK₁ receptor antagonists in combination with selective serotonin reuptake inhibitors in a unit dose of preparation may be varied or adjusted from about 50 mg to about 300 mg of NK₁ receptor antagonists with about 10 mg to about 100 mg of SSRI. An even further quantity of NK₁ receptor
20 antagonists in combination with selective serotonin reuptake inhibitors in a unit dose of preparation may be varied or adjusted from about 50 mg to about 300 mg of NK₁ receptor antagonists with about 20 mg of SSRI, according to the particular application.

The actual dosage employed may be varied depending upon the requirements
25 of the patient and the severity of the condition being treated. Determination of the proper dosage regimen for a particular situation is within the skill of the art. For convenience, the total daily dosage may be divided and administered in portions during the day as required.

The amount and frequency of administration of the compounds of the invention
30 and/or the pharmaceutically acceptable salts thereof will be regulated according to the judgment of the attending clinician considering such factors as age, condition and size of the patient as well as severity of the symptoms being treated. A typical recommended daily dosage regimen for oral administration can range from about 1 mg/day to about 300 mg/day, preferably 1 mg/day to 75 mg/day, in two to four divided
35 doses.

While the present has been described in conjunction with the specific embodiments set forth above, many alternatives, modifications and variations thereof

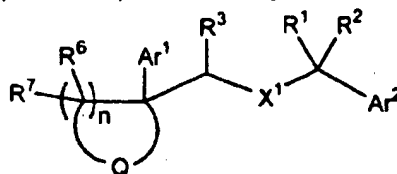
- 100 -

will be apparent to those of ordinary skill in the art. All such alternatives, modifications and variations are intended to fall within the spirit and scope of the present invention.

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We claim:

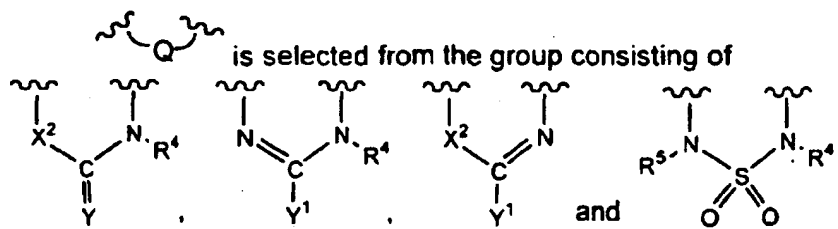
1. A compound represented by the structural formula



- 5 or a pharmaceutically acceptable salt thereof, wherein
 Ar¹ and Ar² are independently selected from the group consisting of R¹⁷-heteroaryl and

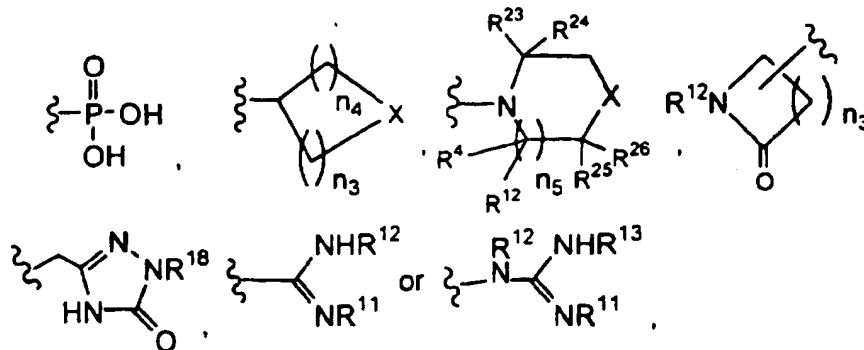


- X¹ is -O-, -S-, -SO-, -SO₂-, -NR¹²-, -N(COR¹²)- or -N(SO₂R¹⁵)-;
 10 R¹, R², R³ and R⁷ are each independently selected from the group consisting of H, C₁-C₆ alkyl, hydroxy(C₁-C₃)alkyl, C₃-C₈ cycloalkyl, -CH₂F, -CHF₂ and -CF₃; or R¹ and R², together with the carbon to which they are attached, form a C₃-C₆ alkylene ring; or, when X¹ is -O-, -S- or -NR¹²-, R¹ and R² together are =O;
 each R⁶ is independently selected from H, C₁-C₆ alkyl, -OR¹³ or -SR¹²;
 15 n is 1-4, if n is greater than 1, then R⁶ and R⁷ can be the same or different on each carbon;



- X² is -O-, -S- or -NR⁵-;
 20 Y is =O, =S or =NR¹¹;
 Y¹ is H, C₁-C₆ alkyl, -NR¹⁷R¹³, -SCH₃, R¹⁹-aryl(CH₂)_{n6}-, R¹⁹-heteroaryl-(CH₂)_{n6}-, -(CH₂)_{n6}-heterocycloalkyl, -(C₁-C₃)alkyl-NH-C(O)O(C₁-C₆)alkyl or -NHC(O)R¹⁵;
 R⁵ is H or -(CH₂)_{n1}-G, wherein n₁ is 0-5, G is H, -CF₃, -CHF₂, -CH₂F, -OH, -O-(C₁-C₆ alkyl), -SO₂R¹³, -O-(C₃-C₈ cycloalkyl), -NR¹³R¹⁴, -SO₂NR¹³R¹⁴, -NR¹³SO₂R¹⁵, -
 25 NR¹³COR¹², -NR¹²(CONR¹³R¹⁴), -CONR¹³R¹⁴, -COOR¹², C₃-C₈ cycloalkyl, R¹⁹-aryl, R¹⁹-heteroaryl,

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or when n_1 is 0, R^5 can also be $-C(O)R^{13}$ or $-C(S)R^{13}$; provided that G is not H when $n_1=0$;

X is $-NR^{20}$, $-N(CONR^{13}R^{14})$, $-N(CO_2R^{13})$, $-N(SO_2R^{15})$, $-N(COR^{12})$,

5 $N(SO_2NHR^{13})$, $-O-$, $-S-$, $-SO-$, $-SO_2-$, $-CF_2-$, $-CH_2-$ or $-CR^{12}F-$;

R^8 , R^9 and R^{10} are independently selected from the group consisting of H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, $-OR^{12}$, halogen, $-CN$, $-NO_2$, $-CF_3$, $-CHF_2$, $-CH_2F$, $-OCF_3$, $-OCHF_2$, $-OCH_2F$, $-COOR^{12}$, $-CONR^{21}R^{22}$, $-NR^{21}COR^{12}$, $-NR^{21}CO_2R^{15}$, $-NR^{21}CONR^{21}R^{22}$, $-NR^{21}SO_2R^{15}$, $-NR^{21}R^{22}$, $-SO_2NR^{21}R^{22}$, $-S(O)_{n_5}R^{15}$, R^{16} -aryl and R^{19} -heteroaryl;

10 R^{11} is H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, $-NO_2$, $-CN$, OH, $-OR^{12}$, $-O(CH_2)_{n_6}R^{12}$, $-(C_1-C_3)alkyl-C(O)NHR^{12}$, R^{19} -aryl $(CH_2)_{n_6}-$ or R^{19} -heteroaryl $-(CH_2)_{n_6}-$;

R^4 and R^{12} are each independently selected from the group consisting of H, C_1 - C_6 alkyl and C_3 - C_8 cycloalkyl;

R^{13} and R^{14} are independently selected from the group consisting of H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, R^{19} -aryl $(CH_2)_{n_6}-$ or R^{19} -heteroaryl $-(CH_2)_{n_6}-$; or R^{13} and R^{14} together are C_3 - C_6 alkylene and with the nitrogen to which they are attached form a 4-7 membered ring, or one of the carbon atoms in the alkylene chain formed by R^{13} and R^{14} is replaced by a heteroatom selected from the group consisting of $-O-$, $-S-$ and $-NR^{12}-$;

20 R^{15} is C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl or $-CF_3$;

R^{16} is 1 to 3 substituents independently selected from the group consisting of H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, C_1 - C_6 alkoxy, halogen and $-CF_3$;

R^{17} is H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, $-COOR^{12}$, $-CONR^{21}R^{22}$, $-NR^{21}R^{22}$, $-NR^{21}COR^{12}$, $-NR^{21}CO_2R^{12}$, $-NR^{21}CONR^{21}R^{22}$, $-NR^{21}SO_2R^{15}$ or $-S(O)_{n_5}R^{15}$;

25 R^{18} is H, C_1 - C_6 alkyl or $-P(O)(OH)_2$;

R^{19} is 1 to 3 substituents independently selected from the group consisting of H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl, $-OH$, halogen, $-CN$, $-NO_2$, $-CF_3$, $-CHF_2$, $-CH_2F$, $-OCF_3$, $-OCHF_2$, $-OCH_2F$, $-O(C_1-C_6 alkyl)$, $-O(C_3-C_8 cycloalkyl)$, $-COOR^{12}$, $-CONR^{21}R^{22}$, $-NR^{21}R^{22}$, $-NR^{21}COR^{12}$, $-NR^{21}CO_2R^{12}$, $-NR^{21}CONR^{21}R^{22}$, $-NR^{21}SO_2R^{15}$ and $-S(O)_{n_5}R^{15}$;

30 R^{20} is H, C_1 - C_6 alkyl, C_3 - C_8 cycloalkyl or $-(CH_2)_{n_6}$ -heterocycloalkyl;

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R^{21} and R^{22} are independently selected from the group consisting of H, C_1-C_6 alkyl, C_3-C_6 cycloalkyl and benzyl; or R^{21} and R^{22} together are C_3-C_6 alkylene and with the nitrogen to which they are attached form a 4-7 membered ring, or one of the carbon atoms in the alkylene chain formed by R^{21} and R^{22} is replaced by a

- 5 heteroatom selected from the group consisting of -O-, -S- and -NR¹²-;

R^{23} , R^{24} , R^{25} and R^{26} are H, C_1-C_6 alkyl and can be together as =O; when $n_3=0$, and R^{25} and $R^{26}=H$, X is not O, N, S;

n_3 and n_4 are independently 1-5, provided that the sum of n_3 and n_4 is 2-6;

n_5 is independently 0-2;

- 10 n_6 is independently 0-3; and

q and r are independently 1 or 2.

2. The compound of claim 1 wherein R^1 , R^3 , R^4 and R^7 are each hydrogen.

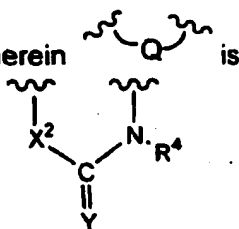
- 15 3. The compound of claim 1 wherein R^6 is H or OH.

4. The compound of claim 1 wherein X^1 is -O- or -NR¹²-.

5. The compound of claim 1 wherein Ar^1 and Ar^2 are each R^8, R^9, R^{10} -phenyl.

20

6. The compound of claim 1 wherein



7. The compound of claim 6 wherein n is 1 or 2.

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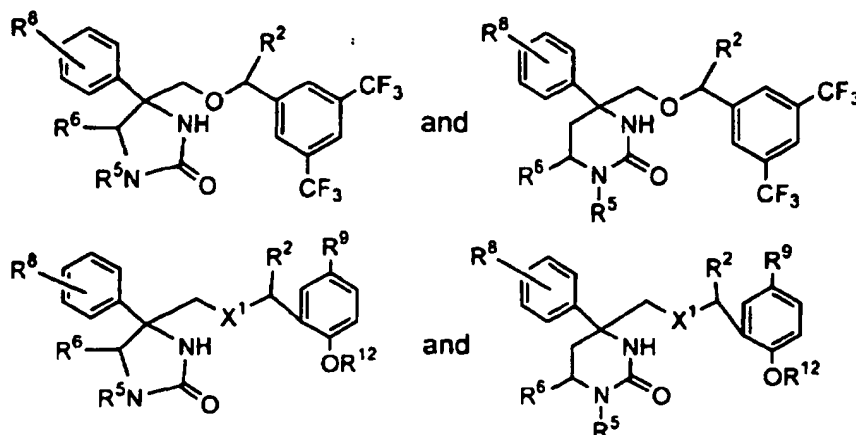
8. The compound of claim 7 wherein X^2 is -NR⁵- and Y is =O.

9. The compound of claim 1 wherein R^1 , R^3 , R^4 and R^7 are each hydrogen; X^1 is -O- or -NR¹²-; Ar^1 and Ar^2 are each R^8, R^9, R^{10} -phenyl; n is 1 or 2; X^2 is -NR⁵-; and

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10. The compound of claim 1 selected from the group of compounds represented by the formula

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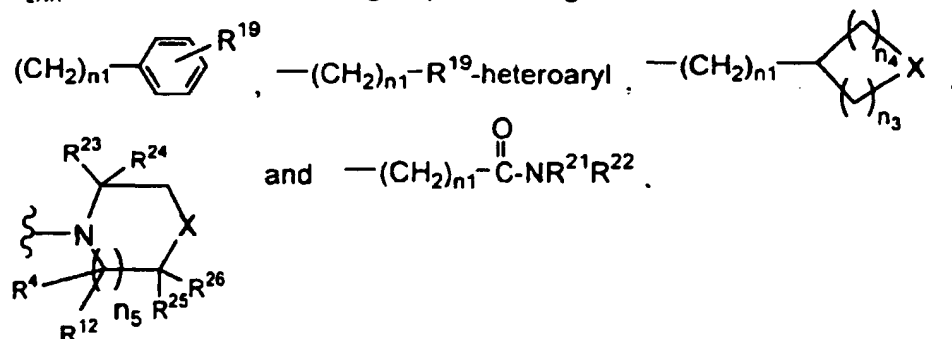


wherein R^8 is H or halogen; R^2 is H, $-\text{CH}_3$ or $-\text{CH}_2\text{OH}$; R^6 is H or $-\text{OH}$; and R^5 is selected from the group consisting of hydrogen and groups of the formula $-(\text{CH}_2)_{n1}-\text{G}$.

5

11. The compound of claim 10 wherein X^1 is $-\text{O}-$, $-\text{NH}-$, $-\text{N}(\text{CH}_3)-$ or $-\text{N}(\text{COCH}_3)-$; R^8 is H or halogen; R^2 is H, $-\text{CH}_3$ or $-\text{CH}_2\text{OH}$; R^6 is H or $-\text{OH}$; R^9 is $-\text{OCF}_3$ or 5-(trifluoromethyl)-1H-tetrazol-1-yl; R^{12} is $-\text{CH}_3$ or cyclopropyl; and R^5 is selected from the group consisting of hydrogen and groups of the formula $-(\text{CH}_2)_{n1}-\text{G}$, wherein $-(\text{CH}_2)_{n1}-\text{G}$ is selected from the group consisting of

10



or $-(\text{CH}_2)_{n1}-\text{G}'$, wherein $n1$ is 2-4 and G' is H, $-\text{OH}$, $-\text{OCH}_3$, ethoxy, isopropoxy, $-\text{O}$ -cyclopropyl, or $-\text{CONR}^{13}\text{R}^{14}$, wherein R^{13} and R^{14} are independently selected from the group consisting of H, $-\text{CH}_3$, ethyl, isopropyl, or cyclopropyl.

15

12. A compound of any of examples 1 to 166.

13. The compound of claim 1 selected from the compounds of Examples: 2, 17, 19, 31, 59, 61, 65, 67, 76, 77, 78, 79a, 79b, 92, 93, 99, 100, 103, 104, 110, 111, 113, 120, 121, 126, 127, 128, 129, 130, 135, 142, 165, 165a, 165b, 166a, 166b or compound 118.

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14. The compound of claim 1 wherein said compound is the compound of Example 61.
- 5 15. The compound of claim 1 wherein said compound is the compound of Example 93.
16. The compound of claim 1 wherein said compound is the compound of Example 2.
- 10 17. The compound of claim 1 wherein said compound is the compound of Example 92.
18. The compound of claim 1 wherein said compound is the compound of Example 127.
- 15 19. The compound of claim 1 wherein said compound is the compound of Example 126.
- 20 20. The compound of claim 1 wherein said compound is the compound of Example 129.
21. The compound of claim 1 wherein said compound is the compound of Example 128.
- 25 22. The compound of claim 1 wherein said compound is the compound of Example 79b.
23. The compound of claim 1 wherein said compound is the compound of Example 79a.
- 30 24. The compound of claim 1 wherein said compound is the compound of Example 166a.
- 35 25. The compound of claim 1 wherein said compound is the compound of Example 166b.

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26. The compound of claim 1 wherein said compound is the compound of Example 165a.
27. The compound of claim 1 wherein said compound is the compound of Example 165b.
28. A pharmaceutical composition comprising an effective amount of a compound of claims 1-27 and a pharmaceutically acceptable carrier.
29. A pharmaceutical composition comprising an effective amount of a compound of claims 1-27, in combination with an effective amount of a selective serotonin reuptake inhibitor and a pharmaceutically acceptable carrier.
30. The use of the compounds of claims 1-27 for the manufacture of a medicament for treating respiratory diseases, inflammatory diseases; skin disorders; ophthalmological disorders; addictions; stress related disorders; obsessive/compulsive disorders; eating disorders; mania; premenstrual syndrome; central nervous system conditions; gastrointestinal disorders; bladder disorders; atherosclerosis; fibrosing disorders; obesity; Type II diabetes; pain related disorders; and genitourinary disorders.
31. The use of the compounds of claims 1-27 in combination with a selective serotonin reuptake inhibitor for the manufacture of a medicament for treating respiratory diseases; inflammatory diseases; skin disorders; ophthalmological disorders; addictions; stress related disorders; obsessive/compulsive disorders; eating disorders; mania; premenstrual syndrome; central nervous system conditions; gastrointestinal disorders; bladder disorders; atherosclerosis; fibrosing disorders; obesity; Type II diabetes; pain related disorders; and genitourinary disorders.
32. The use of the compounds of claims 1-27 for the manufacture of a medicament for treating emesis, depression, anxiety and cough.
33. The use of the compounds of claims 1-27 in combination with a selective serotonin reuptake inhibitor for the manufacture of a medicament for treating treating emesis, depression, anxiety and cough.

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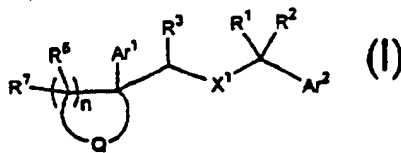
(75) Inventors/Applicants (for US only): SHIH, Neng-Yang [US/US]; 1 Maple Drive, North Caldwell, NJ 07006 (US). SHUE, Ho-Jane [US/US]; 8 Hamilton Place, Pine Brook, NJ 07058 (US). REICHARD, Gregory, A. [US/US]; 4 Bahama Road, Morris Plains, NJ 07950 (US). PALIWAL, Sunil [IN/US]; 245 Spruce Mill Lane, Scotch Plains, NJ 07076 (US). BLYTHIN, David, J. [US/US]; 487 Mountain Avenue, North Caldwell, NJ 07006 (US). PIWINSKI, John, J. [US/US]; 6 Saddle Ridge Drive, Clinton Township, NJ 08833 (US). XIAO, Dong [CN/US]; 124 Spruce Mill Lane, Scotch Plains, NJ 07076 (US). CHEN, Xiao [US/US]; 15 Hillwood Avenue, Edison, NJ 08820 (US).

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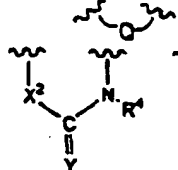
(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, HR, HU, ID, IL, IN, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LU, LV, MA, MD, MG, MK, MN, MX, MZ, NO, NZ, PL, PT, RO, RU.

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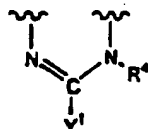
(54) Title: SELECTIVE NEUROKININ ANTAGONISTS



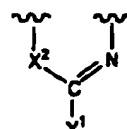
(a)



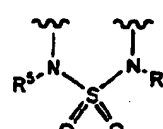
(b)



(c)



(d)



(e)

(57) Abstract: Compound represented by structural formula (I) or a pharmaceutically acceptable salt thereof, wherein Ar¹ and Ar² are optionally substituted heteroaryl or optionally substituted phenyl; X¹ is -O-, -S-, -SO-, -SO₂-, -NR¹²-, -NCOR¹²-, or -NR¹²SO₂R¹⁵; (a) is selected from the group consisting of (b), (c), (d) and (e); X² is -O-, -S- or -NR³-; Y is =O, =S or =NR¹¹; Y¹ is H, C₁-C₆ alkyl, -NR¹⁷R¹⁸, -SCH₃, R¹⁹-aryl(CH₂)_n-, R¹⁹-heteroaryl-(CH₂)_n-, -(CH₂)_n-heterocycloalkyl, -(C₁-C₃)alkyl-NH-C(O)O(C₁-C₆)alkyl or -NHC(O)R¹⁵; R¹, R², R³ and R⁷ are H, alkyl, cycloalkyl, -CHF₂-, -CH₂F or -CF₃; or R¹ and R², together with the carbon to which they are attached, form an alkylene ring; or R¹ and R² together are =O; R⁶ is R⁷ or -OH; and the remaining variables are as defined in the specification, methods of treating diseases susceptible to treatment with neurokinin antagonists with said compounds. Also disclosed are pharmaceutical compositions comprising an effective amount of a compound of claim 1, at least one pharmaceutically acceptable carrier, and in combination with an effective amount of a selective serotonin reuptake inhibitor.

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International Application No
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A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C07D233/32 A61K31/4166 C07D233/22 C07D233/26 C07D233/38 C07D233/46 C07D233/48 C07D233/52 C07D233/42 C07D233/40 C07D233/78 C07D239/10 C07D263/24 C07D285/06 C07D401/04		
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Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
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C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No
A	HARRISON, T. ET AL: "Gem-disubstituted amino-ether based substance P antagonists" BIOORG. MED. CHEM. LETT. (1994), 4(23), 2733-4 XP000995889 the whole document	1-33
A	WO 94 10165 A (MERCK SHARP AND DOHME LTD., UK) 11 May 1994 (1994-05-11) the whole document	1-33
A	EP 0 790 248 A (PFIZER LTD ;PFIZER RES & DEV (IE)) 20 August 1997 (1997-08-20) the whole document	1-33
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Name and mailing address of the ISA European Patent Office, P B 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl. Fax. (+31-70) 340-3016		Authorized officer <div style="text-align: center; font-weight: bold;">Schmid, J-C</div>

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A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D401/12 C07D403/04 C07D403/12 C07D405/04 C07D413/04
 C07D413/10 C07D409/06 C07D521/00 C07F9/22 A61K31/433
 A61K31/513 A61K31/66 A61P1/08 A61P11/06 A61P25/06

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C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No
A	OH, CHANG HYUN ET AL: "Synthesis of new hydantoin-3-acetic acid derivatives" BULL. KOREAN CHEM. SOC. (1988), 9(4), 231-5, XP000995888 table 1	1
A	KNABE, JOACHIM ET AL: "Racemates and enantiomers of basic substituted 5-phenylhydantoins. Syntheses and antiarrhythmic activity" PHARMAZIE (1997), 52(12), 912-919, XP000995887 see compound 4a	1



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A	SCHULTE, KARL E. ET AL: "Hydantoin derivatives as potential antiinflammatory substances" EUR. J. MED. CHEM. - CHIM. THER. (1978), 13(1), 25-31 , XP000995882 tables 1-3	1
P, A	WU, X. ET AL: "Generation of Cyclopenta'c!piperidines and Pyrrolo'3,4-c!piperidines- Potential Substance P Antagonists-from Adducts of Cyclic Dienophiles and 5-Chloro-6-methyl-3-phenyl-2H-1,4-oxazin-2-one" TETRAHEDRON (2000), 56(34), 6279-6290 , XP004214987 the whole document	1-33

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Schmid, J-C

2

INTERNATIONAL SEARCH REPORT

Information on patent family members

National Application No

PCT/US 00/33831

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9410165	A	11-05-1994	AT 188472 T	15-01-2000
			AU 678409 B	29-05-1997
			AU 5342994 A	24-05-1994
			DE 69327541 D	10-02-2000
			DE 69327541 T	10-08-2000
			EP 0666856 A	16-08-1995
			ES 2141174 T	16-03-2000
			JP 8502510 T	19-03-1996
			US 5620989 A	15-04-1997
EP 0790248	A	20-08-1997	CA 2195924 A	28-07-1997
			JP 3026945 B	27-03-2000
			JP 9208582 A	12-08-1997
			US 5846965 A	08-12-1998

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